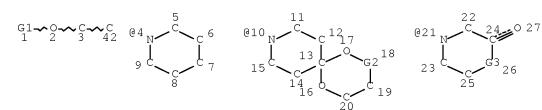
=> d que 167

1 SEA FILE=HCAPLUS ABB=ON PLU=ON US20050215691/PN

L5 STR



VAR G1=4/10/21/34/37

REP G2 = (0-1) CH2

VAR G3=O/N

NODE ATTRIBUTES:

NSPEC IS RC AT 42 NSPEC IS RC AT 43 DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 43

STEREO ATTRIBUTES: NONE

L9 STR

VAR G1=4/10/21/34/37

REP G2=(0-1) CH2

VAR G3=O/N

VAR G4=46/40 NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

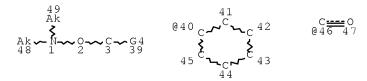
RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 47

STEREO ATTRIBUTES: NONE

L11 7221 SEA FILE=REGISTRY SSS FUL L9

L14 694 SEA FILE=REGISTRY SUB=L11 SSS FUL L5

L17 STR



VAR G4=46/40 NODE ATTRIBUTES: CONNECT IS E1 RC AT 48 CONNECT IS E1 RC AT 49 DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 14

STEREO ATTRIBUTES: NONE

L19 240 SEA FILE=REGISTRY SUB=L11 SSS FUL L17

L21 STR

$$G_1 \sim 0 \sim C \sim G_3 = 0$$
 $G_1 \sim 0 \sim C \sim G_3 = 0$ 
 $G_1 \sim 0 \sim G_3 \sim G_3 = 0$ 
 $G_1 \sim 0 \sim G_3 \sim G_3 = 0$ 
 $G_1 \sim 0 \sim G_3 \sim G_3 \sim G_3 = 0$ 
 $G_1 \sim 0 \sim G_3 \sim$ 

VAR G1=4/10/21/34 REP G2=(0-1) CH2 VAR G3=O/N VAR G4=46/40

NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

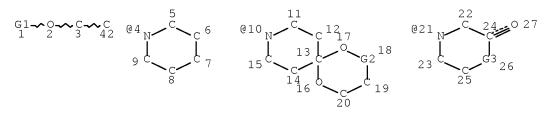
# GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 44

### STEREO ATTRIBUTES: NONE

L23	2091	SEA FILE=REGISTRY SUB=L11 SSS FUL L21
L24	262	SEA FILE=HCAPLUS ABB=ON PLU=ON L14
L25	144	SEA FILE=HCAPLUS ABB=ON PLU=ON L19
L29	209	SEA FILE=HCAPLUS ABB=ON PLU=ON L24(L)PREP/RL
L32	74	SEA FILE=HCAPLUS ABB=ON PLU=ON L29 AND (PLASTIC? OR
		POLYMER?)/SC,SX
L33	60	SEA FILE=HCAPLUS ABB=ON PLU=ON L32 AND (INITIAT? OR
		CATALYST? OR ACTIVAT?)
L34	58	SEA FILE=HCAPLUS ABB=ON PLU=ON L33 NOT L19
L35		STR





VAR G1=4/10/21/34

REP G2=(0-1) CH2

VAR G3=O/N

NODE ATTRIBUTES:

NSPEC IS RC AT 42
NSPEC IS RC AT 43
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

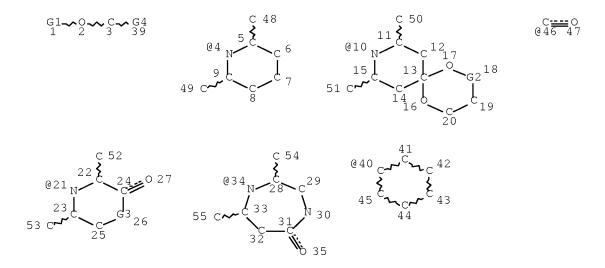
# GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 40

# STEREO ATTRIBUTES: NONE

L37	198	SEA	FILE=REGISTR	Y SUB=L1	4 SSS FUL	ь ьз5	1	
L38	96	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L37		
L39	40	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L34	AND	L38
L42		STR						



VAR G1=4/10/21/34

REP G2=(0-1) CH2

VAR G3=O/N

VAR G4=46/40

NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

### GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 52

STEREO ATTRIBUTES: NONE

DILLICHO	111 11(110011	DD: NONE		
L44	1677	SEA FILE=REGISTRY	SUB=L23 SSS FUI	L42
L45	614	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L44
L46	399	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L45(L)PREP/RL
L47	271	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L46 AND (PLASTIC? OR
		POLYMER?)/SC,SX		
L48	208	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L47 AND (INITIAT? OR
		CATALYST? OR ACTIV	AT?)	
L49	1	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L48 AND L1
L50	2	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L48 AND CLAY?
L51	4	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L48 AND POF/RL
L52	174	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L48 AND CAT/RL
L53	6	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L52 AND MOA/RL
L54	15	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L52 AND (PEP OR PYP)/RL
L55	22	SEA FILE=HCAPLUS A	BB=ON PLU=ON	(L49 OR L50 OR L51) OR
		(L53 OR L54)		
L56	13	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L25 AND L46
L57	35	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L55 OR L56
L58	6	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L39 AND ALKOXYAMINE
		INITIATOR?		
L59	40	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L39 OR L58
L60	5	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L57 AND ALKOXYAMINE
		INITIATOR?		
L61	35	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L57 OR L60
L62	26	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L45 AND ALKOXYAMINE
		INITIATOR?		
L63	26	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L62 AND (PLASTIC? OR
				, , , , , , , , , , , , , , , , , , , ,

POLYMER?)/SC,SX

L64	56	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L61	OR L63
L65	6	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L38	AND ALKOXYAMINE
		INI	TIATOR?				
L66	40	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L59	OR L65
L67	10	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L64	AND L66

#### => d 167 1-10 ibib ed abs hitstr hitind

L67 ANSWER 1 OF 10 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2007:1437481 HCAPLUS Full-text

DOCUMENT NUMBER: 148:215389

TITLE: Synthesis of star polymers via nitroxide mediated

free-radical polymerization: a "core-first"

approach using resorcinarene-based

alkoxyamine initiators

AUTHOR(S): Abraham, Sinoj; Choi, Jae Ho; Ha, Chang-Sik; Kim,

Ιl

CORPORATE SOURCE: Department of Polymer Science and Engineering,

Pusan National University, Pusan, 609-735, S.

Korea

SOURCE: Journal of Polymer Science, Part A: Polymer

Chemistry (2007), 45(23), 5559-5572

CODEN: JPACEC; ISSN: 0887-624X

PUBLISHER: John Wiley & Sons, Inc.

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 19 Dec 2007

AB The synthesis of new octa-functional alkomyamine initiators for nitroxidemediated radical polymerization (NMRP), by the derivatization of resorcinarene
with nitroxide free radicals viz TEMPO and a freshly prepared phosphonylated
nitroxide, is described. The efficiency of these initiators toward the
controlled radical polymerization of styrene and tert-Bu acrylate is studied
in detail. Linear analogs of these multifunctional initiators were also
prepared to compare and evaluate their initiation efficiency. The favorable
conditions for polymerization were optimized by varying the concentration of
initiators and free nitroxides, reaction conditions, etc., to obtain welldefined star polymers. Star polystyrene thus obtained were further used as
macro-initiator for the block copolymn. with tert-Bu acrylate.

IT 1004992-66-8P 1004992-67-9P 1004992-68-0P

1004992-69-1P

(synthesis of octa-functional alkoxyamine

initiators for nitroxide mediated radical polymerization of star

polymers)

RN 1004992-66-8 HCAPLUS

CN INDEX NAME NOT YET ASSIGNED

RN 1004992-67-9 HCAPLUS

CN INDEX NAME NOT YET ASSIGNED

PAGE 1-A

PAGE 2-A

PAGE 3-A

PAGE 4-A

RN 1004992-68-0 HCAPLUS CN INDEX NAME NOT YET ASSIGNED

RN 1004992-69-1 HCAPLUS CN INDEX NAME NOT YET ASSIGNED

PAGE 1-A

- CC 35-4 (Chemistry of Synthetic High Polymers)
- IT Amines, preparation

(alkoxylated, octa-; synthesis of octa-functional alkoxyamine initiators for nitroxide mediated radical polymerization of star polymers)

IT Polymerization

(block, radical; synthesis of octa-armed star polymers using octa-functional alkoxyamine initiators)

IT Nitroxides

(octa-; synthesis of octa-functional alkoxyamine initiators for nitroxide mediated radical polymerization of star polymers)

IT Molecular weight

Polydispersity

(of star polymers prepared using octa-functional alkoxyamine initiators before and after hydrolysis)

IT Polymerization

(radical; synthesis of octa-armed star polymers using octa-functional alkoxyamine initiators)

IT Polymerization catalysts

(radical; synthesis of octa-functional alkoxyamine initiators for nitroxide mediated radical polymerization of star polymers)

IT Polymers, preparation

(star-branched, octa-armed; synthesis of octa-armed star polymers
using octa-functional alkoxyamine initiators)

IT 9003-53-6DP, Polystyrene, hydrolyzed 127972-36-5P, tert-Butyl acrylate-styrene block copolymer

(octa-armed star-shaped; synthesis of octa-armed star polymers using octa-functional alkoxyamine initiators)

IT 366-18-7, 2,2'-Bipyridine 7787-70-4, Cuprous bromide (synthesis of octa-functional alkoxyamine initiators for nitroxide mediated radical polymerization of star polymers)

IT 1004992-66-8P 1004992-67-9P 1004992-68-0P 1004992-69-1P

(synthesis of octa-functional alkoxyamine initiators for nitroxide mediated radical polymerization of star polymers)

TT 75-07-0, Acetaldehyde, reactions 75-64-9, tert-Butylamine, reactions 108-46-3, Resorcinol, reactions 563-76-8, 2-Bromopropionyl bromide 598-02-7, Diethyl phosphate 2564-83-2, TEMPO 65338-98-9 (synthesis of octa-functional alkoxyamine initiators for nitroxide mediated radical polymerization of star polymers)

IT 96183-43-6P 122081-89-4P 1004992-64-6P 1004992-65-7P (synthesis of octa-functional alkoxyamine initiators for nitroxide mediated radical polymerization of star polymers)

REFERENCE COUNT: 83 THERE ARE 83 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L67 ANSWER 2 OF 10 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2005:1311691 HCAPLUS Full-text

DOCUMENT NUMBER: 144:52058

TITLE: Alkoxyamines containing a radically polymerizable

group

INVENTOR(S): Nesvadba, Peter; Kramer, Andreas; Bugnon, Lucienne

PATENT ASSIGNEE(S): Ciba Specialty Chemicals Holding Inc., Switz.

SOURCE: PCT Int. Appl., 54 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.					KIND		DATE			APPLICATION NO.						DATE		
WO	2005118651			A1		20051215		,	WO 2005-EP52260						20050517			
	W:	ΑE,	AG,	AL,	AM,	ΑT,	ΑU,	ΑZ,	BA,	BB,	BG,	BR,	BW,	BY,	BZ,	CA,		
		CH,	CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,		
		GB,	GD,	GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KM,		
		KP,	KR,	KΖ,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,		
		MW,	MX,	MZ,	NA,	NG,	NΙ,	NO,	NΖ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,		
		SC,	SD,	SE,	SG,	SK,	SL,	SM,	SY,	ΤJ,	TM,	TN,	TR,	TT,	TZ,	UA,		
		UG,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	ZW								
	RW:	BW,	GH,	GM,	KE,	LS,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,		
		AM,	AZ,	BY,	KG,	KΖ,	MD,	RU,	ТJ,	TM,	AT,	BE,	BG,	CH,	CY,	CZ,		
		DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	IE,	IS,	IT,	LT,	LU,	MC,		

NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG EP 1749032 Α1 20070207 EP 2005-742775 20050517 EP 1749032 В1 20080227 R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR CN 1957001 Α 20070502 CN 2005-80016626 20050517 JP 2008500307 Τ 20080110 JP 2007-513909 20050517 US 2007232768 A1 20071004 US 2006-596436 20061114 KR 2007024655 Α 20070302 KR 2006-727402 20061227 PRIORITY APPLN. INFO.: EP 2004-102337 A 20040527 WO 2005-EP52260 W 20050517

OTHER SOURCE(S): MARPAT 144:52058

ED Entered STN: 16 Dec 2005

AB The instant invention relates to alkoxyamine initiators/regulators containing an ethylenically unsatd., radically polymerizable group. The compds. are useful for the preparation of complex polymeric architectures. Further aspects of the invention are a polymerizable composition and a polymerization process comprising the alkoxyamine initiators/regulators, a macroinitiator obtainable by the polymerization process and a process for polymerizing with the macroinitiator.

IT 871205-74-2P 871205-75-3P 871205-76-4P 871205-77-5P 871205-78-6P 871205-79-7P 871205-81-1P 871205-82-2P 871205-83-3P 871205-84-4P 871205-85-5P 871205-86-6P 871205-88-8P 871205-89-9P 871205-91-3P 871205-92-4P

(alkoxyamines containing a radically polymerizable group)

RN 871205-74-2 HCAPLUS

CN 2-Propenoic acid, 2-[2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]-1-oxopropoxy]ethyl ester (CA INDEX NAME)

RN 871205-75-3 HCAPLUS

CN 2-Propenoic acid, 2-[2-[(2,6-diethyl-2,3,6-trimethyl-4-oxo-1-piperidinyl)oxy]-1-oxopropoxy]ethyl ester (CA INDEX NAME)

RN 871205-76-4 HCAPLUS

CN 2-Propenoic acid, 2-[[2-[(2,6-diethyl-2,3,6-trimethyl-4-oxo-1-piperidinyl)oxy]-2-methyl-1-oxopropyl]amino]ethyl ester (CA INDEX NAME)

RN 871205-77-5 HCAPLUS

CN 2-Propenoic acid, 1,6-hexanediylbis[imino(1-methyl-2-oxo-2,1-ethanediyl)oxy(2,6-diethyl-2,3,6-trimethyl-1,4-piperidinediyl)] ester (9CI) (CA INDEX NAME)

**∽**Me

RN 871205-78-6 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 2-[[2-[(2,6-diethyl-2,3,6-trimethyl-4-oxo-1-piperidinyl)oxy]-1-oxopropyl]amino]ethyl ester (CA INDEX NAME)

RN 871205-79-7 HCAPLUS

CN 2-Propenoic acid, 2-[[2-[[4-(1,1-dimethylethyl)-2,2-diethyl-6,6-dimethyl-3-oxo-1-piperazinyl]oxy]-1-oxopropyl]amino]ethyl ester (CA INDEX NAME)

RN 871205-81-1 HCAPLUS

CN 2-Propenoic acid, 2,6-diethyl-2,3,6-trimethyl-1-[1-methyl-2-oxo-2-[[2-[(1-oxo-2-propenyl)oxy]ethyl]amino]ethoxy]-4-piperidinyl ester (9CI) (CA INDEX NAME)

RN 871205-82-2 HCAPLUS

CN 1,4-Benzenedicarboxylic acid, bis[2,6-diethyl-2,3,6-trimethyl-1-[1-methyl-2-oxo-2-[[2-[(1-oxo-2-propenyl)oxy]ethyl]amino]ethoxy]-4-piperidinyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 871205-83-3 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 2-[2-[(2,6-diethyl-2,3,6-trimethyl-4-oxo-1-piperidinyl)oxy]-1-oxopropoxy]ethyl ester (CA INDEX NAME)

$$\underbrace{ \text{Et} \quad \text{Me} \quad \text{O-CH-C-O-CH}_2 \text{CH}_2 \text{CH}_2 \text{O-CH}_2 \text{Me} }_{\text{Me}}$$

RN 871205-84-4 HCAPLUS

CN 2-Propenoic acid, 2-methyl-, 2,6-diethyl-2,3,6-trimethyl-1-[1-methyl-2-oxo-2-[2-[(1-oxo-2-propenyl)oxy]ethoxy]ethoxy]-4-piperidinyl ester (9CI) (CA INDEX NAME)

RN 871205-85-5 HCAPLUS

CN 2-Propenoic acid, 2,6-diethyl-2,3,6-trimethyl-1-[1-methyl-2-oxo-2-[2-[(1-oxo-2-propenyl)oxy]ethoxy]ethoxy]-4-piperidinyl ester (9CI) (CA INDEX NAME)

RN 871205-86-6 HCAPLUS

CN 2-Propenoic acid, [[2-[(2,6-diethyl-2,3,6-trimethyl-4-oxo-1-piperidinyl)oxy]-1-oxopropyl]imino]di-2,1-ethanediyl ester (9CI) (CA INDEX NAME)

RN 871205-88-8 HCAPLUS

CN Propanamide, 2-[(2,6-diethyl-2,3,6-trimethyl-4-oxo-1-piperidinyl) oxy]-N-(2-hydroxyethyl)- (CA INDEX NAME)

RN 871205-89-9 HCAPLUS

CN Propanamide, 2-[[4-(1,1-dimethylethyl)-2,2-diethyl-6,6-dimethyl-3-oxo-1-piperazinyl]oxy]-N-(2-hydroxyethyl)- (CA INDEX NAME)

RN 871205-91-3 HCAPLUS

CN Propanamide, 2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]-N-(2-hydroxyethyl)- (CA INDEX NAME)

RN 871205-92-4 HCAPLUS

CN 1,4-Benzenedicarboxylic acid, bis[2,6-diethyl-1-[2-[(2-hydroxyethyl)amino]-1-methyl-2-oxoethoxy]-2,3,6-trimethyl-4-

piperidinyl] ester (9CI) (CA INDEX NAME)

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PAGE 1-B

IT 485844-70-0

(alkoxyamines containing a radically polymerizable group)

RN 485844-70-0 HCAPLUS

CN Propanamide, N,N'-1,6-hexanediylbis[2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]- (CA INDEX NAME)

IC ICM C08F004-00

ICS C07D211-94; C07D241-08; C07F009-40

CC 35-2 (Chemistry of Synthetic High Polymers)

IT 42275-81-0P 639809-62-4P 756490-05-8P 871205-74-2P

871205-75-3P 871205-76-4P 871205-77-5P

871205-78-6P 871205-79-7P 871205-80-0P

871205-81-1P 871205-82-2P 871205-83-3P

871205-84-4P 871205-85-5P 871205-86-6P

871205-88-8P 871205-89-9P 871205-90-2P

871205-91-3P 871205-92-4P

(alkoxyamines containing a radically polymerizable group)

IT 100-20-9, Terephthaloylchloride 141-43-5, Ethanolamine, reactions 814-68-6, Acryloylchloride 2736-37-0, Isobutyrylbromide 3030-47-5

17639-93-9, 2-Chloropropionic acid methylester 51210-48-1 61746-17-6 188065-73-8 188526-94-5 264279-93-8

485844-70-0 871205-87-7

(alkoxyamines containing a radically polymerizable group)

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L67 ANSWER 3 OF 10 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2005:616482 HCAPLUS Full-text

DOCUMENT NUMBER: 144:254433

TITLE: Practical synthesis of alkoxyamine

initiators for living radical

polymerization

AUTHOR(S): Moon, Bongjin; Kang, Minhyuk

CORPORATE SOURCE: Department of Chemistry, Sogang University, Seoul,

121-742, S. Korea

SOURCE: Macromolecular Research (2005), 13(3), 229-235

CODEN: MRAECT; ISSN: 1598-5032

PUBLISHER: Polymer Society of Korea

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 18 Jul 2005

Various alkoxyamine initiators for nitroxide mediated radical polymerization (NMRP) were prepared in high yields by a simple substitution reaction of nitroxide anions with benzyl bromide. The required nitroxide anions were easily generated by treating either nitroxide free radicals or hydroxyamine with an alkali metal such as sodium or potassium in THF. This method is both practical and efficient, since the ionic conditions prevent other side reactions from occurring, such as the self-coupling or oligomerization reactions that are observed in the case of radical trapping conditions. To demonstrate the utility of the resulting alkoxyamine initiators, end- and telechelic-alkoxyamine PEG macroinitiators derived from the alkoxyamines were synthesized by a simple chemical modification, and used for the preparation of PEG-b-PS and PS-b-PEG-b-PS block copolymers by NMRP.

IT 877141-19-0P 877141-21-4P

(macroinitiator; synthesis of alkoxyamine
initiators for living radical polymerization and block copolymer
made by using them)

RN 877141-19-0 HCAPLUS

CN Poly(oxy-1,2-ethanediyl),  $\alpha$ -[1,4-dioxo-4-[2-phenyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethoxy]butyl]- $\omega$ -methoxy- (9CI) (CA INDEX NAME)

RN 877141-21-4 HCAPLUS

CN Poly(oxy-1,2-ethanediyl),  $\alpha$ -[1,4-dioxo-4-[2-phenyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethoxy]butyl]- $\omega$ -[1,4-dioxo-4-[2-phenyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethoxy]butoxy]- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

$$-CH_2-CH_2-C-O-CH_2-CH-O-N$$
Me
Me
Me
Me
Me

IT 154554-67-3P, 2,2,6,6-Tetramethyl-1-(1-phenylethoxy)piperidine 161776-41-6P

(synthesis of alkoxyamine initiators for living

radical polymerization and block copolymer made by using them)

RN 154554-67-3 HCAPLUS

CN Piperidine, 2,2,6,6-tetramethyl-1-(1-phenylethoxy)- (CA INDEX NAME)

RN 161776-41-6 HCAPLUS

CN Benzeneethanol,  $\beta$ -[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]- (CA INDEX NAME)

IT 877141-11-2P

(synthesis of alkoxyamine initiators for living radical polymerization and block copolymer made by using them)

RN 877141-11-2 HCAPLUS

CN Piperidine, 1-[2-[[(1,1-dimethylethyl)dimethylsilyl]oxy]-1-phenylethoxy]-2,2,6,6-tetramethyl- (CA INDEX NAME)

CC 35-3 (Chemistry of Synthetic High Polymers)

ST alkoxyamine living radical polymn initiator synthesis; substitution reaction living radical polymn catalyst synthesis; polyethylene glycol macroinitiator alkoxyamine block copolymer manuf; nitroxide mediated radical polymn catalyst synthesis

IT Polyoxyalkylenes, reactions

(synthesis of alkoxyamine initiators for living

radical polymerization and block copolymer made by using them)

IT 877141-19-0P 877141-21-4P

(macroinitiator; synthesis of alkoxyamine

initiators for living radical polymerization and block copolymer made by using them)

IT 154554-67-3P, 2,2,6,6-Tetramethyl-1-(1-phenylethoxy)piperidine 161776-41-6P 227000-59-1P 727989-07-3P

(synthesis of alkoxyamine initiators for living

radical polymerization and block copolymer made by using them)

IT 108-30-5, Succinic anhydride, reactions 585-71-7,

(1-Bromoethyl) benzene 2564-83-2, TEMPO 7440-09-7, Potassium, reactions 7440-23-5, Sodium, reactions 9004-74-4, Polyethylene

glycol monomethyl ether 25322-68-3, Polyethylene glycol 41252-83-9, 2-Bromo-2-phenylethanol 61015-94-9 98194-56-0,

tert-Butylchlorosilane

(synthesis of alkoxyamine initiators for living

radical polymerization and block copolymer made by using them)

IT 31961-02-1P 37684-51-8P 245759-95-9P 877141-11-2P 877141-14-5P

(synthesis of alkoxyamine initiators for living

radical polymerization and block copolymer made by using them)

IT 697291-44-4P, Ethylene oxide-styrene diblock copolymer 697303-36-9P, Ethylene oxide-styrene triblock copolymer

(synthesis of alkozyamine initiators for living

radical polymerization and block copolymer made by using them)

REFERENCE COUNT: 28 THERE ARE 28 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L67 ANSWER 4 OF 10 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2005:157125 HCAPLUS Full-text

DOCUMENT NUMBER: 142:392795

TITLE: Construction of a polymer skeleton that is cut in

half by ionizing radiation

AUTHOR(S): Shimizu, Takashi; Ichikawa, Tsuneki

CORPORATE SOURCE: Division of Molecular Chemistry, Graduate School

of Engineering, Hokkaido University, Sapporo,

060-8628, Japan

SOURCE: Journal of Polymer Science, Part A: Polymer

Chemistry (2005), 43(5), 1068-1075 CODEN: JPACEC; ISSN: 0887-624X

PUBLISHER: John Wiley & Sons, Inc.

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 24 Feb 2005

AΒ Polystyrene with a benzyl ester of carboxylic acid at the center of a polymer skeleton was synthesized by living radical polymerization. The initiator used had two functional groups for 2,2,6,6-tetramethylpiperidinoxyl (TEMPO)mediated living radical polymerization on the benzyl and the carboxylic sides of the benzyl ester. Introduction of the benzyl ester changed the polystyrene from a crosslink type to a scission type polymer on  $\gamma$ -irradiation Irradiation of the polymer resulted in a binary change of the mol. weight because of the dissociative capture of secondary electrons by the benzyl ester, as:  $MnR1COOCH(C6H5)R2Mn + e \rightarrow MnR1COO + CH(C6H5)R2Mn$ . The binary change of the mol. weight suggests that the polymer can be used as a new type of radiation resist with high sensitivity and spatial resolution to ionizing and high resistivity to plasma etching. The number of scissions per 100 eV radiation energy absorbed was 0.29, which was about one fourth of the yield of secondary electrons. The low efficiency was because of the recombination of polymer radicals generated by the dissociative electron attachment. ΙT 647849-32-9P

(construction of a benzyl ester central functionalized polystyrene skeleton that is cut in half by ionizing radiation)

RN 647849-32-9 HCAPLUS

IT 209550-24-3P

(construction of a benzyl ester central functionalized polystyrene skeleton that is cut in half by ionizing radiation)

RN 209550-24-3 HCAPLUS

CN Benzenemethanol, 4-[1-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl]-(CA INDEX NAME)

CC 35-8 (Chemistry of Synthetic High Polymers)

IT Polymerization

Polymerization catalysts

(living, radical; construction of a benzyl ester central functionalized polystyrene skeleton that is cut in half by ionizing radiation)

IT 647849-32-9P

(construction of a benzyl ester central functionalized polystyrene skeleton that is cut in half by ionizing radiation)

IT 209550-24-3P

(construction of a benzyl ester central functionalized polystyrene skeleton that is cut in half by ionizing radiation)

REFERENCE COUNT: 25 THERE ARE 25 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L67 ANSWER 5 OF 10 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2004:2855 HCAPLUS  $\underline{\text{Full-text}}$ 

DOCUMENT NUMBER: 140:77932

TITLE: Cationic alkoxyamines and their use in producing

nanoparticles from natural or synthetic

clays

INVENTOR(S): Muehlebach, Andreas; Nesvadba, Peter; Kramer,

Andreas

PATENT ASSIGNEE(S): Ciba Specialty Chemicals Holding Inc., Switz.

SOURCE: PCT Int. Appl., 62 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.					KIN	D	DATE	TE APPLICATION NO.						DATE			
WO	2004000809					_	2003	20031231			2003-	20030617					
	W:	ΑE,	AG,	AL,	AM,	ΑT,	ΑU,	AZ,	BA,	BB,	BG,	BR,	BY,	BZ,	CA,	CH,	
		CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	
		GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	ΚE,	KG,	KP,	KR,	KZ,	
		LC,	LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	
		ΝI,	NO,	NZ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	
		SL,	ТJ,	TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US,	UZ,	VC,	VN,	YU,	ZA,	
		ZM,	ZW														
	RW:	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AM,	AZ,	
		BY,	KG,	KΖ,	MD,	RU,	ТJ,	TM,	ΑT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	
		EE,	ES,	FI,	FR,	GB,	GR,	HU,	ΙE,	ΙΤ,	LU,	MC,	NL,	PT,	RO,	SE,	
		SI,	SK,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	$\mathrm{ML}_{m{\prime}}$	MR,	
		ΝE,	SN,	TD,	ΤG												
CA	2486	958			A1		2003	1231		CA 2	2003-	2486	958		2	0030617	
AU	2003	2793	73		A1 20040106				AU 2003-279373						20030617		
EP	1515	950			A1	A1 20050323			EP 2003-740262					20030617			
	R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	ΙΤ,	LI,	LU,	NL,	SE,	MC,	
		PT,	IE,	SI,	LT,	LV,	FΙ,	RO,	MK,	CY,	AL,	TR,	BG,	CZ,	EE,	HU, SK	
	1662						2005	0831	CN 2003-814665						2	0030617	
JP	2005	5389	64		T 20051222			1222	JP 2004-514745					20030617			
MX	2004	PA12	885		Α		2005	0331		MX 2	2004-	PA12	885		2	0041217	
US	2005	2156	91		A1		2005	0929		US 2	2004-	5190	30		2	0041222	
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ORIT	Y APP	LN.	INFO	.:						EP 2	2002-	4055	20		A 2	0020624	
										WO 2	2003-	EP63	70	,	W 2	0030617	

OTHER SOURCE(S): MARPAT 140:77932

ED Entered STN: 02 Jan 2004

GΙ

$$\begin{array}{c} \text{Me} \\ \text{Me} \\ \text{Me} \\ \text{Me} \\ \text{Me} \\ \text{Et} \\ \text{Et} \\ \text{Et} \\ \text{I} \end{array}$$

The instant invention relates to cationic alkoxyamines such as I, which are useful as polymerization initiators/regulators in a controlled stable free radical polymerization of unsatd. compds. in the presence of nanoparticles of natural or synthetic clays to produce intercalated and/or exfoliated nanoparticles. The invention also relates to improved nanocomposites produced by this process and to the use of these nanocomposite compns. as, for example, coatings, sealants, caulks, adhesives and as plastic additives.

II 639809-49-7P, 1-tert-Butyl-4-[1-[4-

(chloromethyl)phenyl]ethoxy]-3,3-diethyl-5,5-dimethylpiperazin-2-one 639809-51-1P, 1-tert-Butyl-3,3-diethyl-5,5-dimethyl4-[1-[4-(4-methylpiperazin-1-ylcarbonyl)phenyl]ethoxy]piperazin-2-one 639809-54-4P, 2-(2,6-Diethyl-2,3,6-trimethylpiperidin-1-yloxy)-N-(3-dimethylaminopropyl)propionamide 639809-56-6P, 2-(2,6-Diethyl-4-hydroxy-2,3,6-trimethylpiperidin-1-yloxy)-N-(3-dimethylaminopropyl)propionamide 639809-60-2P, 2,6-Diethyl-1-(1-phenylethoxy)-2,3,6-trimethylpiperidin-4-one 0-(3-dimethylaminopropyl) oxime 639809-63-5P, Bis[2,6-diethyl-1-[1-(3-dimethylaminopropylcarbamoyl)ethoxy]-2,3,6-trimethylpiperidin-4-yl] terephthalate 639809-65-7P, N-(3-Dimethylaminopropyl)-2-(4-hydroxy-2,2,6,6-tetramethylpiperidin-1-yloxy)propionamide 639809-67-9P, 2-(2,6-Diethyl-4-hydroxy-2,3,6-trimethylpiperidin-1-yloxy)-N-(3-dimethylaminopropyl)-2-methylpropionamide 639809-73-7P

(catalyst precursor; cationic alkoxyamines for catalysts/regulators for polymerization of unsatd. compds. in presence of nanoparticles from natural or synthetic clays for manufacture of nanocomposites)

RN 639809-49-7 HCAPLUS

CN Piperazinone, 4-[1-[4-(chloromethyl)phenyl]ethoxy]-1-(1,1-dimethylethyl)-3,3-diethyl-5,5-dimethyl- (9CI) (CA INDEX NAME)

RN 639809-51-1 HCAPLUS

CN Piperazine, 1-[4-[1-[[4-(1,1-dimethylethyl)-2,2-diethyl-6,6-dimethyl-3-oxo-1-piperazinyl]oxy]ethyl]benzoyl]-4-methyl- (9CI) (CA INDEX NAME)

RN 639809-54-4 HCAPLUS

CN Propanamide, 2-[(2,6-diethyl-2,3,6-trimethyl-1-piperidinyl)oxy]-N-[3-(dimethylamino)propyl]- (CA INDEX NAME)

RN 639809-56-6 HCAPLUS

CN Propanamide, 2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]-N-[3-(dimethylamino)propyl]- (CA INDEX NAME)

RN 639809-60-2 HCAPLUS

CN 4-Piperidinone, 2,6-diethyl-2,3,6-trimethyl-1-(1-phenylethoxy)-, O-[3-(dimethylamino)propyl]oxime (CA INDEX NAME)

RN 639809-63-5 HCAPLUS

CN 1,4-Benzenedicarboxylic acid, bis[1-[2-[[3-(dimethylamino)propyl]amino]-1-methyl-2-oxoethoxy]-2,6-diethyl-2,3,6-trimethyl-4-piperidinyl] ester (9CI) (CA INDEX NAME)

Me<sub>2</sub>N-(CH<sub>2</sub>)<sub>3</sub>-NH-C-CH-O Et Me O Me Et Me O Et Me

PAGE 1-B

RN 639809-65-7 HCAPLUS

CN Propanamide, N-[3-(dimethylamino)propyl]-2-[(4-hydroxy-2,2,6,6-tetramethyl-1-piperidinyl)oxy]- (CA INDEX NAME)

RN 639809-67-9 HCAPLUS

CN Propanamide, 2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]-N-[3-(dimethylamino)propyl]-2-methyl- (CA INDEX NAME)

RN 639809-73-7 HCAPLUS

CN Propanoic acid, 2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]-, 3-bromopropyl ester (CA INDEX NAME)

CN Benzoic acid, 4-[1-[[4-(1,1-dimethylethyl)-2,2-diethyl-6,6-dimethyl-3-oxo-1-piperazinyl]oxy]ethyl]- (CA INDEX NAME)

RN 478697-26-6 HCAPLUS CN 4-Piperidinone, 2,6-diethyl-2,3,6-trimethyl-1-(1-phenylethoxy)-, oxime (CA INDEX NAME)

IT 639809-48-6P, [4-[1-(4-tert-Butyl-2,2-diethyl-6,6-dimethyl-3-oxopiperazin-1-yloxy)ethyl]benzyl]triethylammonium chloride 639809-50-0P, 4-[4-[1-(4-tert-Butyl-2,2-diethyl-6,6-dimethyl-3-oxopiperazin-1-yloxy)ethyl]benzoyl]-1,1-dimethylpiperazin-1-ium iodide 639809-52-2P, [3-[2-(2,6-Diethyl-2,3,6-trimethylpiperidin-1-yloxy)propionylamino]propyl]dimethylethylammonium bromide 639809-55-5P, [3-[2-(2,6-Diethyl-4-hydroxy-2,3,6-trimethylpiperidin-1-yloxy)propionylamino]propyl]dimethylethylammonium bromide 639809-59-9P, [3-[2,6-Diethyl-1-(1-phenylethoxy)-2,3,6-trimethylpiperidin-4-ylideneaminooxy]propyl]dimethylethylammonium

m bromide 639809-61-3F, Bis[[3-[2-(2,6-diethyl-4-hydroxy-2,3,6-trimethylpiperidin-1-yloxy)propionylamino]propyl]dimethylethylam monium bromide] terephthalate 639809-64-6P, Ethyl[3-[2-(4-hydroxy-2,2,6,6-tetramethylpiperidin-1-yloxy)propionylamino]propyl]dimethylammonium bromide 639809-66-8P, [3-[2-(2,6-Diethyl-4-hydroxy-2,3,6-trimethylpiperidin-1-yloxy)-2-methylpropionylamino]propyl]dimethylethy lammonium bromide 639809-69-1P, Benzyl[3-[2-(2,6-Diethyl-4-hydroxy-2,3,6-trimethylpiperidin-1-yloxy)-2-methylpropionylamino]propyl]dimethylammonium chloride 639809-70-4P, Benzyl[3-[2-(2,6-Diethyl-4-hydroxy-2,3,6-trimethylpiperidin-1-yloxy)propionylamino]propyl]dimethylammonium chloride 639809-71-5P, [3-[2-(2,6-Diethyl-4-hydroxy-2,3,6-trimethylpiperidin-1-yloxy)propionyloxy]propyl]tributylphosphonium bromide

(cationic alkoxyamines for catalysts/regulators for polymerization of unsatd. compds. in presence of nanoparticles from natural or synthetic clays for manufacture of nanocomposites)

RN 639809-48-6 HCAPLUS

CN Benzenemethanaminium, 4-[1-[[4-(1,1-dimethylethyl)-2,2-diethyl-6,6-dimethyl-3-oxo-1-piperazinyl]oxy]ethyl]-N,N,N-triethyl-, chloride (9CI) (CA INDEX NAME)

RN 639809-50-0 HCAPLUS

CN Piperazinium, 4-[4-[1-[[4-(1,1-dimethylethyl)-2,2-diethyl-6,6-dimethyl-3-oxo-1-piperazinyl]oxy]ethyl]benzoyl]-1,1-dimethyl-, iodide (9CI) (CA INDEX NAME)

RN 639809-52-2 HCAPLUS

CN 1-Propanaminium, 3-[[2-[(2,6-diethyl-2,3,6-trimethyl-1-piperidinyl)oxy]-1-oxopropyl]amino]-N-ethyl-N,N-dimethyl-, bromide (9CI) (CA INDEX NAME)

Br-

RN 639809-55-5 HCAPLUS

CN 1-Propanaminium, 3-[[2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]-1-oxopropyl]amino]-N-ethyl-N,N-dimethyl-, bromide (9CI) (CA INDEX NAME)

Br-

RN 639809-59-9 HCAPLUS

CN 1-Propanaminium, 3-[[[2,6-diethyl-2,3,6-trimethyl-1-(1-phenylethoxy)-4-piperidinylidene]amino]oxy]-N-ethyl-N,N-dimethyl-, bromide (9CI) (CA INDEX NAME)

● Br-

RN 639809-61-3 HCAPLUS

CN 1-Propanaminium, 3,3'-[1,4-phenylenebis[carbonyloxy(2,6-diethyl-2,3,6-trimethyl-4,1-piperidinediyl)oxy(2-methyl-1-oxo-2,1-ethanediyl)imino]]bis[N-ethyl-N,N-dimethyl-, dibromide (9CI) (CA INDEX NAME)

PAGE 1-A

●2 Br-

PAGE 1-B

RN 639809-64-6 HCAPLUS

CN 1-Propanaminium, N-ethyl-3-[[2-[(4-hydroxy-2,2,6,6-tetramethyl-1-piperidinyl)oxy]-1-oxopropyl]amino]-N,N-dimethyl-, bromide (9CI) (CA INDEX NAME)

Br -

RN 639809-66-8 HCAPLUS

CN 1-Propanaminium, 3-[[2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]-2-methyl-1-oxopropyl]amino]-N-ethyl-N,N-dimethyl-, bromide (9CI) (CA INDEX NAME)

$$\begin{array}{c|c} \text{Et} & \text{Me} & \text{Me} \\ \text{NO} & \text{C-C-NH-(CH2)} \\ \text{Me} & \text{Me} \end{array}$$

● Br-

RN 639809-69-1 HCAPLUS

CN Benzenemethanaminium, N-[3-[[2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]-2-methyl-1-oxopropyl]amino]propyl]-N,N-dimethyl-, chloride (9CI) (CA INDEX NAME)

● c1-

RN 639809-70-4 HCAPLUS

CN Benzenemethanaminium, N-[3-[[2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]-1-oxopropyl]amino]propyl]-N,N-dimethyl-, chloride (9CI) (CA INDEX NAME)

● c1-

RN 639809-71-5 HCAPLUS

CN Phosphonium, tributyl[3-[2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]-1-oxopropoxy]propyl]-, bromide (9CI) (CA INDEX NAME)

Br -

IC ICM C07D211-94

ICS C07D241-08; C07F009-38; C08F002-00

CC 37-3 (Plastics Manufacture and Processing) Section cross-reference(s): 23, 27, 28, 38, 42

ST cationic alkoxyamine initiator unsatd compd polymn nanocomposite manuf; plastic additive nanocomposite cationic alkoxyamine polymn initiator; caulking nanocomposite cationic alkoxyamine polymn initiator; adhesive nanocomposite cationic alkoxyamine polymn initiator; tertiary butyldiethyloxopiperazinyloxyethyl benzyltriethylammonium chloride initiator polymn nanocomposite manuf; sealant nanocomposite cationic alkoxyamine polymn initiator; coating nanocomposite cationic alkoxyamine polymn initiator

IT Phosphonium compounds

Quaternary ammonium compounds, preparation (alkoxyamino; cationic alkoxyamines for catalysts /regulators for polymerization of unsatd. compds. in presence of nanoparticles from natural or synthetic clays for manufacture of nanocomposites)

IT Disperse systems

(cationic alkoxyamines for catalysts/regulators for polymerization of unsatd. compds. in presence of nanoparticles from natural or synthetic clays for manufacture of nanocomposite dispersions)

IT Nanocomposites

Polymerization catalysts

(cationic alkoxyamines for catalysts/regulators for polymerization of unsatd. compds. in presence of nanoparticles from natural or synthetic clays for manufacture of nanocomposites)

IT Phyllosilicate minerals

Smectite-group minerals

(cationic alkoxyamines for catalysts/regulators for polymerization of unsatd. compds. in presence of nanoparticles from natural or synthetic clays for manufacture of nanocomposites)

IT Adhesives

(cationic alkoxyamines for catalysts/regulators for polymerization of unsatd. compds. in presence of nanoparticles from natural or synthetic clays for manufacture of nanocomposites for adhesive additives)

IT Coating materials

(cationic alkoxyamines for catalysts/regulators for polymerization of unsatd. compds. in presence of nanoparticles from natural or synthetic clays for manufacture of nanocomposites for coating additives)

IT Inks

(cationic alkoxyamines for catalysts/regulators for

polymerization of unsatd. compds. in presence of nanoparticles from natural or synthetic clays for manufacture of nanocomposites for ink additives)

IT Paints

(cationic alkoxyamines for catalysts/regulators for polymerization of unsatd. compds. in presence of nanoparticles from natural or synthetic clays for manufacture of nanocomposites for paint additives)

IT Clay minerals

(intercalated, cationic alkoxyamine-; cationic alkoxyamines for catalysts/regulators for polymerization of unsatd. compds. in presence of nanoparticles from natural or synthetic clays for manufacture of nanocomposites)

IT Plastics, miscellaneous

(thermoplastics; cationic alkoxyamines for catalysts /regulators for polymerization of unsatd. compds. in presence of nanoparticles from natural or synthetic clays for manufacture of nanocomposites for thermoplastics)

188526-94-5P 639809-49-7P, 1-tert-Butyl-4-[1-[4-ΙT (chloromethyl)phenyl]ethoxy]-3,3-diethyl-5,5-dimethylpiperazin-2-one 639809-51-1P, 1-tert-Butyl-3,3-diethyl-5,5-dimethyl4-[1-[4-(4methylpiperazin-1-ylcarbonyl)phenyl]ethoxy]piperazin-2-one 639809-53-3P, 2-Chloro-N-(3-dimethylaminopropyl)propionamide 639809-54-4P, 2-(2,6-Diethyl-2,3,6-trimethylpiperidin-1-yloxy)-N-(3-dimethylaminopropyl)propionamide 639809-56-6P, 2-(2,6-Diethyl-4-hydroxy-2,3,6-trimethylpiperidin-1-yloxy)-N-(3dimethylaminopropyl)propionamide 639809-58-8P, Diethyl [1-[tert-butyl-[1-(3-dimethylaminopropylcarbamoyl)ethoxy]amino]-2,2dimethylpropyl]phosphonate 639809-60-2P, 2,6-Diethyl-1-(1-phenylethoxy)-2,3,6-trimethylpiperidin-4-one O-(3-dimethylaminopropyl) oxime 639809-63-5P, Bis [2,6-diethyl-1-[1-(3-dimethylaminopropylcarbamoyl)ethoxy]-2,3,6trimethylpiperidin-4-yl] terephthalate 639809-65-7P, N-(3-Dimethylaminopropyl)-2-(4-hydroxy-2,2,6,6-tetramethylpiperidin-1yloxy)propionamide 639809-67-9P, 2-(2,6-Diethyl-4-hydroxy-2,3,6-trimethylpiperidin-1-yloxy)-N-(3-dimethylaminopropyl)-2methylpropionamide 639809-68-0P, 2-Bromo-N-(3-dimethylaminopropyl)-2methylpropionamide 639809-72-6P, 3-Bromopropyl 2-bromopropionate 639809-73-7P

(catalyst precursor; cationic alkoxyamines for catalysts/regulators for polymerization of unsatd. compds. in presence of nanoparticles from natural or synthetic clays for manufacture of nanocomposites)

74-88-4, Methyl iodide, reactions 74-96-4, Ethvl bromide Benzyl chloride, reactions 109-01-3, N-Methylpiperazine 109-54-6, 3-Dimethylaminopropyl chloride 109-55-7, 3-Dimethylamino-1propylamine 121-44-8, Triethylamine, reactions 627-18-9 998-40-3, Tributylphosphine 1592-20-7, 4-Chloromethylstyrene 2226-96-2, 4-Hydroxy-TEMPO 17639-93-9, Methyl 2-chloropropionate 20769-85-1, 2-Bromo-2-methylpropionyl bromide 61745-37-7, 2,6-Diethyl-4-hydroxy-2,3,6-trimethylpiperidine 1-N-oxyl 61746-17-6, 2,6-Diethyl-2,3,6-trimethylpiperidine 1-N-oxyl 264279-93-8, 1-tert-Butyl-3,3-diethyl-5,5-dimethylpiperazin-2-one 4-N-oxyl 319458-08-7, 4-[1-(4-tert-Butyl-2,2-diethyl-6,6-dimethyl-3oxopiperazin-1-yloxy)ethyl]benzoic acid 478697-26-6 639809-62-4

(catalyst precursor; cationic alkoxyamines for catalysts/regulators for polymerization of unsatd. compds. in presence of nanoparticles from natural or synthetic clays for manufacture of nanocomposites)

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ΙT
     639809-48-6P, [4-[1-(4-tert-Butyl-2,2-diethyl-6,6-dimethyl-3-
     oxopiperazin-1-yloxy)ethyl]benzyl]triethylammonium chloride
     639809-50-0P, 4-[4-[1-(4-tert-Butyl-2,2-diethyl-6,6-dimethyl-3-
     oxopiperazin-1-yloxy)ethyl]benzoyl]-1,1-dimethylpiperazin-1-ium iodide
     639809-52-2P, [3-[2-(2,6-Diethyl-2,3,6-trimethylpiperidin-1-
     vloxy)propionylamino|propyl|dimethylethylammonium bromide
     639809-55-5P, [3-[2-(2,6-Diethyl-4-hydroxy-2,3,6-
     trimethylpiperidin-1-yloxy)propionylamino]propyl]dimethylethylammonium
               639809-57-7P 639809-59-9P, [3-[2,6-Diethyl-1-(1-
     phenylethoxy)-2,3,6-trimethylpiperidin-4-ylideneaminooxy]propyl]dimeth
     ylethylammonium bromide 639809-61-3P, Bis[[3-[2-(2,6-diethyl-
     4-hydroxy-2,3,6-trimethylpiperidin-1-yloxy)propionylamino]propyl]dimet
     hylethylammonium bromide] terephthalate 639809-64-6P,
     Ethyl[3-[2-(4-hydroxy-2,2,6,6-tetramethylpiperidin-1-
     yloxy)propionylamino]propyl]dimethylammonium bromide
     639809-66-8P, [3-[2-(2,6-Diethyl-4-hydroxy-2,3,6-
     trimethylpiperidin-1-yloxy)-2-methylpropionylamino]propyl]dimethylethy
     lammonium bromide 639809-69-1P, Benzyl[3-[2-(2,6-Diethyl-4-
     hydroxy-2,3,6-trimethylpiperidin-1-yloxy)-2-
     methylpropionylamino]propyl]dimethylammonium chloride
     639809-70-4P, Benzyl[3-[2-(2,6-Diethyl-4-hydroxy-2,3,6-
     trimethylpiperidin-1-yloxy)propionylamino]propyl]dimethylammonium
     chloride 639809-71-5P, [3-[2-(2,6-Diethyl-4-hydroxy-2,3,6-
     trimethylpiperidin-1-yloxy)propionyloxy]propyl]tributylphosphonium
     bromide
        (cationic alkoxyamines for catalysts/regulators for
       polymerization of unsatd. compds. in presence of nanoparticles from
       natural or synthetic clays for manufacture of nanocomposites)
     9003-49-0P, Poly(butyl acrylate) 9003-53-6P, Polystyrene
ΙT
        (cationic alkoxyamines for catalysts/regulators for
       polymerization of unsatd. compds. in presence of nanoparticles from
        natural or synthetic clays for manufacture of nanocomposites)
ΙT
     1318-00-9, Vermiculite
                             1318-74-7, Kaolinite, uses 1318-93-0D,
     Montmorillonite, derivs. 1319-41-1, Saponite 12172-85-9,
     Beidellite 12173-47-6, Optigel SH 12174-06-0, Nontronite
     (Fe2(Si3.67Al0.33)Na0.33(OH)2O10.xH2O) 12244-16-5, Endellite
     12417-86-6, Stevensite
                             565450-32-0, Nanofil EXM588
        (cationic alkoxyamines for catalysts/regulators for
        polymerization of unsatd. compds. in presence of nanoparticles from
       natural or synthetic clays for manufacture of nanocomposites)
                               THERE ARE 2 CITED REFERENCES AVAILABLE FOR
REFERENCE COUNT:
                               THIS RECORD. ALL CITATIONS AVAILABLE IN THE
                               RE FORMAT
L67 ANSWER 6 OF 10 HCAPLUS COPYRIGHT 2008 ACS on STN
                         2003:583946 HCAPLUS Full-text
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         139:246241
TITLE:
                         An improved catalytic method for alkoxyamine
                         synthesis - functionalized and biradical
                         initiators for nitroxide-mediated radical
                        polymerization
AUTHOR(S):
                        Bothe, Marc; Schmidt-Naake, Gudrun
CORPORATE SOURCE:
                        Institut fuer Technische Chemie, Technische
                         Universitaet Clausthal, Clausthal-Zellerfeld,
                         38678, Germany
SOURCE:
                        Macromolecular Rapid Communications (2003),
                         24(10), 609-613
                        CODEN: MRCOE3; ISSN: 1022-1336
                        Wiley-VCH Verlag GmbH & Co. KGaA
PUBLISHER:
DOCUMENT TYPE:
                        Journal
```

LANGUAGE: English

ED Entered STN: 30 Jul 2003

AB Mn(salen)Cl [(N,N'-disalicylidene-1,2-ethanediamino)manganese chloride] was applied as a low-cost catalyst for the formation of alkoxyamines from nitroxides and substituted styrenes. These "unimol. initiators" for nitroxide-mediated radical polymerization (NMRP) were synthesized using 2,2,6,6-tetramethyl-1-piperidine- 1-oxyl and 2,2,5-trimethyl-4-phenyl-3-azahexane-3-oxyl. Functionalized alkoxyamines were obtained from 4-vinylbenzyl chloride and 4-vinylbenzyl alc. The divinyl compound 1,2-bis(4-vinylphenyl)ethane was converted to an alkoxyamine monomer and to bisaminooxy compds., which can be used as "biradical initiators" for NMRP.

IT 154554-67-3P 212132-38-2P 596135-22-7P 596135-24-9P

(improved catalytic method for synthesis of alkoxyamine functionalized and biradical initiators for nitroxide-mediated radical polymerization)

RN 154554-67-3 HCAPLUS

CN Piperidine, 2,2,6,6-tetramethyl-1-(1-phenylethoxy)- (CA INDEX NAME)

RN 212132-38-2 HCAPLUS

CN Piperidine, 1-[1-[4-(chloromethyl)phenyl]ethoxy]-2,2,6,6-tetramethyl-(CA INDEX NAME)

RN 596135-22-7 HCAPLUS

CN Piperidine, 1,1'-[1,2-ethanediylbis(4,1-phenyleneethylideneoxy)]bis[2,2,6,6-tetramethyl-(9CI) (CA INDEX NAME)

RN 596135-24-9 HCAPLUS

CN Benzenemethanamine, N,N'-[1,2-ethanediylbis(4,1phenyleneethylideneoxy)]bis[N-(1,1-dimethylethyl)- $\alpha$ -(1methylethyl) - (9CI) (CA INDEX NAME)

596135-23-8P ΙT

> (monomer; improved catalytic method for synthesis of alkoxyamine functionalized and biradical initiators for nitroxide-mediated radical polymerization)

596135-23-8 HCAPLUS RN

Piperidine, 1-[1-[4-[2-(4-ethenylphenyl)ethyl]phenyl]ethoxy]-2,2,6,6-CN tetramethyl- (CA INDEX NAME)

- CC 35-3 (Chemistry of Synthetic High Polymers)
- ST alkoxyamine initiator nitroxide radical polymn
- Catalysts ΙT

(for synthesis of alkoxyamine functionalized and biradical initiators for nitroxide-mediated radical polymerization)

Polymerization catalysts ΤТ

> (radical; improved catalytic method for synthesis of alkoxyamine functionalized and biradical initiators for nitroxide-mediated radical polymerization)

53177-12-1P ΙT

> (catalyst; improved catalytic method for synthesis of alkoxyamine functionalized and biradical initiators for nitroxide-mediated radical polymerization)

154554-67-3P 212132-38-2P 227000-59-1P ΙT

227000-85-3P 433266-98-9P 596135-22-7P

596135-24-9P

(improved catalytic method for synthesis of alkoxyamine functionalized and biradical initiators for nitroxide-mediated radical polymerization)

1074-61-9P, 4-Vinylbenzyl alcohol 2628-16-2P, 4-Vinylphenyl acetate ΙT 48174-52-3P, 1,2-Bis(4-vinylphenyl)ethane

> (intermediate; improved catalytic method for synthesis of alkoxyamine functionalized and biradical initiators for nitroxide-mediated radical polymerization)

ΤT 596135-23-8P

(monomer; improved catalytic method for synthesis of alkoxyamine

functionalized and biradical initiators for nitroxide-mediated radical polymerization)

IT 94-93-9, N,N'-Disalicylidene-1,2-ethanediamine 1592-20-7,

4-Vinylbenzyl chloride 2564-83-2, TEMPO 6156-78-1, Manganese

acetate tetrahydrate 61015-94-9

(starting material; improved catalytic method for synthesis of alkoxyamine functionalized and biradical initiators for

nitroxide-mediated radical polymerization)

REFERENCE COUNT: 30 THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L67 ANSWER 7 OF 10 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2003:383971 HCAPLUS <u>Full-text</u>

DOCUMENT NUMBER: 139:117728

TITLE: Synthesis and reactivity of functionalized

alkoxyamine initiators for

nitroxide-mediated radical polymerization of

styrene

AUTHOR(S): Li, Irene Q.; Knauss, Daniel M.; Priddy, Duane B.;

Howell, Bob A.

CORPORATE SOURCE: Department of Chemistry and Geochemistry, Colorado

School of Mines, Golden, CO, 80401, USA

SOURCE: Polymer International (2003), 52(5), 805-812

CODEN: PLYIEI; ISSN: 0959-8103

PUBLISHER: John Wiley & Sons Ltd.

DOCUMENT TYPE: Journal LANGUAGE: English

ED Entered STN: 20 May 2003

The synthesis and examination of different functionalized (2,2,6,6-AΒ tetramethyl-1-piperidinyloxy free radical) TEMPO-containing alkoxyamine initiators for nitroxide-mediated radical polymerization of styrene are reported. Initiators with ester and carbonate functional groups were synthesized by a low-temperature radical-abstraction reaction of the functionalized ethylbenzene in the presence of TEMPO to introduce the functional groups onto the initiating chain-end of polystyrene. An initiator with two alkoxyamine groups sym. located at each end of a carbonate bond was also synthesized and used for nitroxide-mediated styrene polymerization Styrene polymerization using these initiators followed first-order kinetics up to approx. 60 min at 140° or 30% monomer conversion. Alkoxyamines bearing an acetoxy or tert-butylcarbonate group at the p-position of 1-(2,2,6,6tetramethyl- 1-piperidinyloxy)ethylbenzene behave in a similar way to the unfunctionalized initiator. With an initiator containing two alkoxyamine groups, the resulting polymer mol. weight was twice that of the polymer obtained from initiators with only one alkoxyamine group, as expected from propagation from both chain-ends. Upon hydrolysis of the carbonate bond, equivalent polymer chain growth occurred from each alkoxyamine site in the difunctional initiator.

IT 213699-59-3P 224824-56-0P 562102-19-6P 562102-23-2P

(synthesis of TEMPO-containing alkoxyamine initiators for radical polymerization of styrene)

RN 213699-59-3 HCAPLUS

CN Phenol, 4-[1-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl]-, acetate (ester) (9CI) (CA INDEX NAME)

224824-56-0 HCAPLUS RN

Carbonic acid, 1,1-dimethylethyl 4-[1-[(2,2,6,6)-tetramethyl-1-CN piperidinyl)oxy]ethyl]phenyl ester (9CI) (CA INDEX NAME)

562102-19-6 HCAPLUS RN

CN Phenol, 4-[(1,1-dimethylethoxy)] [(2,2,6,6-tetramethyl-1piperidinyl)oxy]methyl]-, acetate (ester) (9CI) (CA INDEX NAME)

562102-23-2 HCAPLUS RN

CN Phenol, 4-[1-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl]-,carbonate (2:1) (ester) (9CI) (CA INDEX NAME)

ΙT 154554-67-3P 562102-22-1P

(synthesis of TEMPO-containing alkoxyamine initiators

for radical polymerization of styrene)

154554-67-3 HCAPLUS RN

Piperidine, 2,2,6,6-tetramethyl-1-(1-phenylethoxy)- (CA INDEX NAME) CN

RN 562102-22-1 HCAPLUS

CN Phenol, 4-[1-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl]- (CA INDEX NAME)

CC 35-4 (Chemistry of Synthetic High Polymers)

ST TEMPO contg alkoxyamine initiator synthesis styrene radical polymn

IT Amines, preparation

(alkoxylated; synthesis of TEMPO-containing alkoxyamine initiators for radical polymerization of styrene)

IT Polydispersity

(of polystyrene prepared using TEMPO-containing alkoxyamine initiators)

IT Polymerization

(radical, solution; of styrene using TEMPO-containing alkoxyamine initiators and synthesis of the initiators)

IT Polymerization kinetics

(radical; of styrene using TEMPO-containing alkoxyamine initiators and synthesis of the initiators)

IT Polymerization datalysts

(radical; synthesis of TEMPO-containing alkoxyamine initiators for radical polymerization of styrene)

IT Nitroxides

(synthesis of TEMPO-containing alkoxyamine initiators for radical polymerization of styrene)

IT 9003-53-6P, Polystyrene

(radical polymerization of styrene using TEMPO-containing alkoxyamine initiators and synthesis of the initiators)

IT 213699-59-3P 224824-56-0P 562102-19-6P 562102-23-2P

(synthesis of TEMPO-containing alkoxyamine initiators for radical polymerization of styrene)

TT 75-44-5, Phosgene 123-07-9, 4-Ethylphenol 1876-22-8, Di-tert-butylperoxyoxalate 2564-83-2, TEMPO 2628-16-2, 4-Acetoxystyrene 224824-55-9

(synthesis of TEMPO-containing alkoxyamine initiators for radical polymerization of styrene)

IT 3245-23-6P, 4-(Acetoxy)ethylbenzene 154554-67-3P
562102-22-1P

(synthesis of TEMPO-containing alkoxyamine initiators

for radical polymerization of styrene)

REFERENCE COUNT: 74 THERE ARE 74 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L67 ANSWER 8 OF 10 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2001:247383 HCAPLUS Full-text

DOCUMENT NUMBER: 134:281264

TITLE: Polymerization compositions using

nitrogen-containing free radicals

INVENTOR(S): Lai, John Ta-yuan; Filla, Deborah S.

PATENT ASSIGNEE(S): The B.F. Goodrich Company, USA

SOURCE: PCT Int. Appl., 39 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PA:	PATENT NO.					KIND DATE			APPLICATION NO.						DATE			
WO	WO 2001023435												20000929					
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		CN,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EE,	ES,	FI,	GB,	GD,	GE,	GH,		
		GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KΕ,	KG,	KP,	KR,	KΖ,	LC,	LK,		
		LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NO,	NΖ,		
		PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	ΤJ,	TM,	TR,	TT,	TΖ,		
		UA,	UG,	UZ,	VN,	YU,	ZA,	ZW,	AM,	ΑZ,	BY,	KG,	KΖ,	MD,	RU,	ΤJ,	TM	
	RW:	GH,	GM,	KE,	LS,	MW,	MΖ,	SD,	SL,	SZ,	TZ,	UG,	ZW,	AT,	BE,	CH,		
		CY,	DE,	DK,	ES,	FI,	FR,	GB,	GR,	ΙE,	ΙΤ,	LU,	MC,	NL,	PT,	SE,		
		BF,	ΒJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GW,	$\mathrm{ML}_{\prime}$	MR,	ΝE,	SN,	TD,	ΤG		
EP	1259	551			A1 20021127		EP 2000-968537			37		2	00009	929				
	R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	ΙΤ,	LI,	LU,	NL,	SE,	MC,		
		PT,	ΙE,	SI,	LT,	LV,	FΙ,	RO,	MK,	CY,	AL							
PRIORIT	Y APP	LN.	INFO	.:					1	US 1	999-	4093.	29		A 1	99909	930	
									1	US 2	000-	1722	19P		P 2	0000	110	
									1	US 1	999-:	1722	19P	:	P 1	99909	930	
									1	WO 2	000-1	JS27	038	1	w 2	00009	929	

OTHER SOURCE(S): MARPAT 134:281264

ED Entered STN: 06 Apr 2001

AB Nitrogen-containing free radicals and free radical precursors are used to effect efficient, controlled polymerization of polymeric materials, including monomers, to form polymers, including homopolymers, copolymers, and block polymers. In particular, nitroxide compds., methods of making nitroxide compds., methods of using the nitroxide compds. to polymerize monomer compns., and polymer compns. made using the nitroxide compds. are disclosed. Thus, heating tetramethylmorpholone 8.64 with p-xylene 2.66 and Mo oxide 0.2 at reflux, dropwisely adding tert-Bu hydroperoxide (I) 16 over 1 h, refluxing for 20 min, further adding I 20 g while heating for another 1 h and working up gave an adduct useful for radical precursor.

IT 332380-43-5P 332380-44-6P

(radical polymerization initiator precursor; polymerization compns. using nitrogen-containing free radicals)

RN 332380-43-5 HCAPLUS

CN 2-Morpholinone, 4,4'-[1,4-phenylenebis(methyleneoxy)]bis[3,3,5,5-

tetramethyl- (9CI) (CA INDEX NAME)

RN 332380-44-6 HCAPLUS

CN 2-Morpholinone, 3,3,5,5-tetramethyl-4-(1-phenylethoxy)- (CA INDEX NAME)

IC ICM C08F004-00

ICS C08F002-38; C08F293-00; C07D265-32; C07D241-08; C07C239-20

CC 35-3 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 67

ST morpholone xylene adduct radical polymn initiator

IT Polymerization catalysts

(polymerization compns. using nitrogen-containing free radicals)

IT 1313-27-5, Molybdenum oxide, uses

(nitroxide forming catalysts; polymerization compns. using nitrogen-containing free radicals)

IT 332380-43-5P 332380-44-6P

(radical polymerization initiator precursor; polymerization compns. using nitrogen-containing free radicals)

IT 113872-32-5P

(radical polymerization initiator; polymerization compns. using nitrogen-containing free radicals)

IT 100-41-4, Ethylbenzene, reactions 106-42-3, p-Xylene, reactions 90032-83-0, 3,3,5,5-Tetramethyl-2-morpholinone

(reactant for polymerization initiator precursor; polymerization compns. using nitrogen-containing free radicals)

REFERENCE COUNT:

THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L67 ANSWER 9 OF 10 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2001:229230 HCAPLUS <u>Full-text</u>

DOCUMENT NUMBER: 135:33673

TITLE: Thermolysis of Free-Radical Initiators:

tert-Butylazocumene and Its 1,3- and 1,4-Bisazo

and 1,3,5-Trisazo Analogues

AUTHOR(S): Engel, Paul S.; Pan, Li; Ying, Yunming; Alemany,

Lawrence B.

CORPORATE SOURCE: Department of Chemistry, Rice University, Houston,

TX, 77251, USA

SOURCE: Journal of the American Chemical Society (2001),

123(16), 3706-3715

CODEN: JACSAT; ISSN: 0002-7863

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 01 Apr 2001

AB Four tert-butylazocumenes (I-IV) were prepared from the corresponding cyanobenzenes, and their nitrogen evolution kinetics and products were analyzed. In combination with TEMPO, the simplest compound, tert-butylazocumene (I), shows promise as a "one-radical" initiator of styrene polymerization An ABNO-trapped cumyl radical is a particularly stable trialkylhydroxylamine, whose thermolysis half-life is 2.1 h at 150 °C. Taking advantage of this stability, we trapped the cumyl radical centers from IV to afford tris adduct. While the behavior of the meta bisazoalkane III can be mostly predicted from that of I, the para isomer II exhibits both unusual products and kinetics, attributed to the formation of quinodimethane via azocontaining radical. Finally, evidence is presented that IV is a possible initiator of star polymerization of Me methacrylate.

IT 344299-94-1P 344299-95-2P 344299-99-6P

(thermolysis of the free-radical initiators 1,3- and 1,4-bisazo and 1,3,5-trisazo tert-butylazocumene analogs)

RN 344299-94-1 HCAPLUS

CN Piperidine, 1,1',1''-[1,3,5-benzenetriyltris[(1-methylethylidene)oxy]]tris[2,2,6,6-tetramethyl-(9CI) (CA INDEX NAME)

RN 344299-95-2 HCAPLUS

CN Piperidine, 1,1'-[1,4-phenylenebis[(1-methylethylidene)oxy]]bis[2,2,6,6-tetramethyl- (9CI) (CA INDEX NAME)

```
RN
     344299-99-6 HCAPLUS
CN
     9-Azabicyclo[3.3.1]nonane, 9,9',9''-[1,3,5-benzenetriyltris[(1-
     methylethylidene)oxy]]tris- (9CI) (CA INDEX NAME)
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
     35-3 (Chemistry of Synthetic High Polymers)
     Section cross-reference(s): 21
     kinetics thermolysis butylazocumene radical polymn initiator
ST
    Activation enthalpy
ΤТ
       Activation entropy
     Radical scavengers
        (in relation to thermolysis of the free-radical initiators
        1,3- and 1,4-bisazo and 1,3,5-trisazo tert-butylazocumene analogs)
ΙT
     Activation energy
     Simulation and Modeling, physicochemical
        (of the thermolysis of the free-radical initiators 1,3-
        and 1,4-bisazo and 1,3,5-trisazo tert-butylazocumene analogs)
ΙT
     Polymerization catalysts
        (radical; in relation to thermolysis of the free-radical
        initiators 1,3- and 1,4-bisazo and 1,3,5-trisazo
        tert-butylazocumene analogs)
ΙT
     Thermal decomposition
     Thermal decomposition kinetics
        (thermolysis of the free-radical initiators 1,3- and
        1,4-bisazo and 1,3,5-trisazo tert-butylazocumene analogs)
     40014-21-9P 344299-68-9P 344299-72-5P
ΤТ
                                               344299-76-9P
        (thermolysis of the free-radical initiators 1,3- and
        1,4-bisazo and 1,3,5-trisazo tert-butylazocumene analogs)
ΙT
     2564-83-2, TEMPO
        (thermolysis of the free-radical initiators 1,3- and
        1,4-bisazo and 1,3,5-trisazo tert-butylazocumene analogs)
ΙT
     31785-68-9P, 9-Azabicyclo[3.3.1]non-9-yloxy
        (thermolysis of the free-radical initiators 1,3- and
        1,4-bisazo and 1,3,5-trisazo tert-butylazocumene analogs)
     4148-81-6P
                 7518-53-8P 10365-94-3P, 1,3,5-Tricyanobenzene
TT
     19937-49-6P
                 95627-96-6P 344299-83-8P
                                               344299-85-0P
        (thermolysis of the free-radical initiators 1,3- and
        1,4-bisazo and 1,3,5-trisazo tert-butylazocumene analogs)
     585-32-0P, Cumylamine
ΙT
        (thermolysis of the free-radical initiators 1,3- and
        1,4-bisazo and 1,3,5-trisazo tert-butylazocumene analogs)
TΤ
     98-83-9, reactions
                        108-98-5, Thiophenol, reactions
     2388-14-9
               33581-95-2
        (thermolysis of the free-radical initiators 1,3- and
        1,4-bisazo and 1,3,5-trisazo tert-butylazocumene analogs)
ΙT
     4132-49-4P 4148-93-0P 9003-53-6P, Polystyrene 9011-14-7P, PMMA
     19956-03-7P 344299-87-2P
                                 344299-88-3P 344299-91-8P
     344299-92-9P 344299-93-0P 344299-94-1P
     344299-95-2P
                  344299-96-3P
                                  344299-97-4P
                                                344299-98-5P
     344299-99-6P
        (thermolysis of the free-radical initiators 1,3- and
        1,4-bisazo and 1,3,5-trisazo tert-butylazocumene analogs)
REFERENCE COUNT:
                         94
                               THERE ARE 94 CITED REFERENCES AVAILABLE FOR
                               THIS RECORD. ALL CITATIONS AVAILABLE IN THE
                              RE FORMAT
L67 ANSWER 10 OF 10 HCAPLUS COPYRIGHT 2008 ACS on STN
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1999:558851 HCAPLUS Full-text

ACCESSION NUMBER:

DOCUMENT NUMBER: 132:152244

TITLE: Synthesis and properties of polymeric networks

prepared by "living" free radical polymerization

and end-linking processes

AUTHOR(S): Chaumont, Philippe; Asgarzadeh, Firouz;

Ourdouillie, Pascal; Beyou, Emmanuel; Mechin,

Francoise; Dumon, Michel

CORPORATE SOURCE: Unite Mixte de Recherches "Ingenierie des

Materiaux Macromoleculaires", Universite,

Villeurbanne, 69622, Fr.

SOURCE: Polymer Preprints (American Chemical Society,

Division of Polymer Chemistry) (1999), 40(2),

366-367

CODEN: ACPPAY; ISSN: 0032-3934

American Chemical Society, Division of Polymer PUBLISHER:

Chemistry

DOCUMENT TYPE: Journal LANGUAGE: English

Entered STN: 02 Sep 1999

Polymer networks were synthesized by "living" free radical polymerization, i.e. the free radical synthesis of difunctional precursors, followed by the crosslinking of these precursors. Three types of controlled polymerization were studied to prepare the precursors and the networks: (a) reversible termination with nitroxide type control agents, (b) atom transfer radical polymerization, and (c) radical addition-fragmentation transfer. The structure and the swelling properties of the gels formed were studied.

257955-86-5P ΙT

> (free radical control agent; for living free radical polymerization by reversible termination with nitroxide radicals)

RN 257955-86-5 HCAPLUS

1,4-Benzenedicarboxylic acid, bis[2-phenyl-2-[(2,2,6,6-tetramethyl-1-CN piperidinyl)oxy]ethyl] ester (9CI) (CA INDEX NAME)

35-4 (Chemistry of Synthetic High Polymers) CC

ΙT 257955-86-5P

> (free radical control agent; for living free radical polymerization by reversible termination with nitroxide radicals)

7787-70-4, Copper bromide (CuBr) 71071-44-8, ΙT 78-67-1, AIBN

4,4'-Di-n-heptyl-2,2'-bipyridine

(polymerization catalyst; synthesis and properties of polymeric networks prepared by living free radical polymerization and end-linking processes)

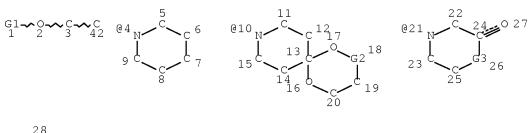
REFERENCE COUNT: 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

=> d que 168

1 SEA FILE=HCAPLUS ABB=ON PLU=ON US20050215691/PN

L5 STR



VAR G1=4/10/21/34/37

REP G2=(0-1) CH2

VAR G3=O/N

NODE ATTRIBUTES:

NSPEC IS RC AT 42 NSPEC IS RC AT 43 DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 43

STEREO ATTRIBUTES: NONE

L9 STR

VAR G1=4/10/21/34/37

VAR G3=O/N VAR G4=46/40 NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

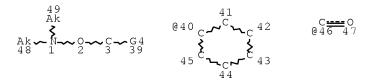
NUMBER OF NODES IS 47

STEREO ATTRIBUTES: NONE

L11 7221 SEA FILE=REGISTRY SSS FUL L9

L14 694 SEA FILE=REGISTRY SUB=L11 SSS FUL L5

L17 STR



VAR G4=46/40
NODE ATTRIBUTES:
CONNECT IS E1 RC AT 48
CONNECT IS E1 RC AT 49
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 14

STEREO ATTRIBUTES: NONE

L19 240 SEA FILE=REGISTRY SUB=L11 SSS FUL L17

L21 STR

VAR G1=4/10/21/34

REP G2=(0-1) CH2

VAR G3=O/N

VAR G4=46/40

#### NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

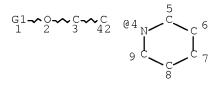
#### GRAPH ATTRIBUTES:

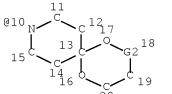
RING(S) ARE ISOLATED OR EMBEDDED

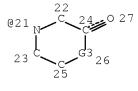
NUMBER OF NODES IS 44

#### STEREO ATTRIBUTES: NONE

L23	2091	SEA FILE=REGISTRY SUB=L11 SSS FUL L21
L24	262	SEA FILE=HCAPLUS ABB=ON PLU=ON L14
L25	144	SEA FILE=HCAPLUS ABB=ON PLU=ON L19
L29	209	SEA FILE=HCAPLUS ABB=ON PLU=ON L24(L)PREP/RL
L32	74	SEA FILE=HCAPLUS ABB=ON PLU=ON L29 AND (PLASTIC? OR
		POLYMER?)/SC,SX
L33	60	SEA FILE=HCAPLUS ABB=ON PLU=ON L32 AND (INITIAT? OR
		CATALYST? OR ACTIVAT?)
L34	58	SEA FILE=HCAPLUS ABB=ON PLU=ON L33 NOT L19
L35		STR







G1-O-C-C 39 40 41 43

VAR G1=4/10/21/34

REP G2=(0-1) CH2

VAR G3=O/N

NODE ATTRIBUTES:

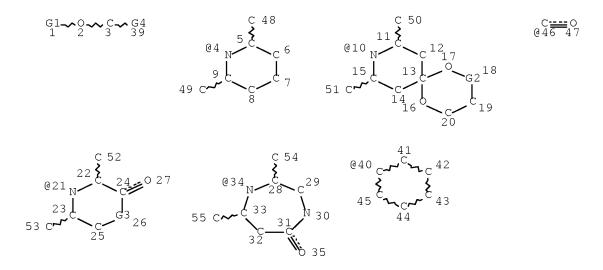
NSPEC IS RC AT 42
NSPEC IS RC AT 43
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

#### GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 40

#### STEREO ATTRIBUTES: NONE

			. • —					
L37	198	SEA	FILE=REGISTR	Y SUB=L1	4 SSS FU	L L35	ō	
L38	96	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L37		
L39	40	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L34	AND	L38
L42		STR						



VAR G1=4/10/21/34

REP G2=(0-1) CH2

VAR G3=O/N

VAR G4=46/40

NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

#### GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 52

#### STEREO ATTRIBUTES: NONE

L44	1677	SEA FILE=REGISTRY SUB=L23 SSS FUL L42
L45	614	SEA FILE=HCAPLUS ABB=ON PLU=ON L44
L46	399	SEA FILE=HCAPLUS ABB=ON PLU=ON L45(L)PREP/RL
L47	271	SEA FILE=HCAPLUS ABB=ON PLU=ON L46 AND (PLASTIC? OR
		POLYMER?)/SC,SX
L48	208	SEA FILE=HCAPLUS ABB=ON PLU=ON L47 AND (INITIAT? OR
		CATALYST? OR ACTIVAT?)
L49	1	SEA FILE=HCAPLUS ABB=ON PLU=ON L48 AND L1
L50	2	SEA FILE=HCAPLUS ABB=ON PLU=ON L48 AND CLAY?
	4	SEA FILE=HCAPLUS ABB=ON PLU=ON L48 AND POF/RL
L52	174	SEA FILE=HCAPLUS ABB=ON PLU=ON L48 AND CAT/RL
L53	6	SEA FILE=HCAPLUS ABB=ON PLU=ON L52 AND MOA/RL
L54	15	SEA FILE=HCAPLUS ABB=ON PLU=ON L52 AND (PEP OR PYP)/RL
L55	22	SEA FILE=HCAPLUS ABB=ON PLU=ON (L49 OR L50 OR L51) OR
		(L53 OR L54)
L56	13	SEA FILE=HCAPLUS ABB=ON PLU=ON L25 AND L46
L57	35	SEA FILE=HCAPLUS ABB=ON PLU=ON L55 OR L56
L58	6	SEA FILE=HCAPLUS ABB=ON PLU=ON L39 AND ALKOXYAMINE
		INITIATOR?
L59	40	SEA FILE=HCAPLUS ABB=ON PLU=ON L39 OR L58
L60	5	SEA FILE=HCAPLUS ABB=ON PLU=ON L57 AND ALKOXYAMINE
		INITIATOR?
L61	35	SEA FILE=HCAPLUS ABB=ON PLU=ON L57 OR L60
L62	26	SEA FILE=HCAPLUS ABB=ON PLU=ON L45 AND ALKOXYAMINE
		INITIATOR?
L63	26	SEA FILE=HCAPLUS ABB=ON PLU=ON L62 AND (PLASTIC? OR
	_	, , , , , , , , , , , , , , , , , , , ,

#### POLYMER?)/SC,SX

L64	56	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L61	OR L63
L65	6	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L38	AND ALKOXYAMINE
		INI	TIATOR?				
L66	40	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L59	OR L65
L67	10	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L64	AND L66
L68	46	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L64	NOT L67

=> sel 168 hit rn 1-E58 THROUGH E256 ASSIGNED

=> d 168 1-46 ibib ed abs fhitstr hitind

L68 ANSWER 1 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2007:772026 HCAPLUS Full-text

DOCUMENT NUMBER: 147:323380

TITLE: Site-selective surface-initiated

polymerization by Langmuir-Blodgett lithography AUTHOR(S): Brinks, Marion K.; Hirtz, Michael; Chi, Lifeng;

Fuchs, Harald; Studer, Armido

CORPORATE SOURCE: Organisch-Chemisches Institut and NRW Graduate

School of Chemistry, Westfaelische

Wilhelms-Universitaet, Muenster, 48149, Germany Angewandte Chemie, International Edition (2007),

46(27), 5231-5233

CODEN: ACIEF5; ISSN: 1433-7851 Wiley-VCH Verlag GmbH & Co. KGaA

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 147:323380

ED Entered STN: 17 Jul 2007

AB Brushing the surface: Structured polymer brushes are readily prepared by site-selective immobilization of initiators in a self-assembly process by Langmuir-Blodgett lithog. with subsequent polymerization. The AFM images show a DPPC/alkoxy amine LB film before and after surface-initiated controlled nitroxide-mediated radical polymerization. Large surface areas (several cm2) can readily be structured with this method.

IT 947770-00-5P

SOURCE:

PUBLISHER:

(synthesis of alkoxy amines and site-selective immobilization thereof by Langmuir-Blodgett lithog. for preparation of polymer brushes)

RN 947770-00-5 HCAPLUS

CN Piperidine, 2,2,6,6-tetramethyl-1-[2-methyl-1-[4-[[[10-(triethoxysilyl)decyl]oxy]methyl]phenyl]propoxy]- (CA INDEX NAME)

Eto-
$$S_{i-}^{\text{OEt}}$$
 (CH<sub>2</sub>)<sub>10</sub>-O-CH<sub>2</sub>  $i-Pr$  Me Ne Me

CC 35-4 (Chemistry of Synthetic High Polymers)

ST initiator lithog nitroxide radical polymn polymer brush prepn

Lithography ΙT Self-assembly (for site-selective immobilization of initiators for preparation of polymer brushes) Polymerization ΙT (graft; preparation of polymer brushes with site-selective immobilization of initiators by Langmuir-Blodgett lithog. and subsequent radical polymerization) Polymer brushes ΙT (preparation of polymer brushes with site-selective immobilization of initiators by Langmuir-Blodgett lithog. and subsequent radical polymerization) Polvmerization ΙT (radical; preparation of polymer brushes with site-selective immobilization of initiators by Langmuir-Blodgett lithog. and subsequent radical polymerization) ΙT Polymerization catalysts (radical; synthesis of alkoxy amines and site-selective immobilization thereof by Langmuir-Blodgett lithog. for preparation of polymer brushes) Polymerization ΙT (surface; preparation of polymer brushes with site-selective immobilization of initiators by Langmuir-Blodgett lithog. and subsequent radical polymerization) 9003-49-0P, Butyl acrylate homopolymer 9003-53-6P, Polystyrene TΤ (grafted on silica layer on silicon wafer; preparation of polymer brushes with site-selective immobilization of initiators by Langmuir-Blodgett lithog. and subsequent radical polymerization) 897033-80-6 154554-67-3 ΙT (preparation of polymer brushes with site-selective immobilization of initiators by Langmuir-Blodgett lithog. and subsequent radical polymerization) 63-89-8, L- $\alpha$ -Dipalmitoylphosphatidylcholine ΤT (preparation of polymer brushes with site-selective immobilization of initiators by Langmuir-Blodgett lithog. and subsequent radical polymerization) 7631-86-9P, Silica, preparation ΙT (surface layer, polystyrene- or poly(Bu acrylate)-grafted; preparation of polymer brushes with site-selective immobilization of initiators by Langmuir-Blodgett lithog. and subsequent radical polymerization) 947770-00-5P 947770-04-9P ΙT (synthesis of alkoxy amines and site-selective immobilization thereof by Langmuir-Blodgett lithog. for preparation of polymer brushes) ΙT 947769-94-0P 947769-95-1P 947769-96-2P 947769-97-3P 947769-98-4P 947769-99-5P 947770-01-6P 947770-02-7P 947770-03-8P (synthesis of alkoxy amines and site-selective immobilization thereof by Langmuir-Blodgett lithog. for preparation of polymer brushes) 7440-21-3, Silicon, miscellaneous ΙT (wafer, substrate; preparation of polymer brushes with site-selective immobilization of initiators by Langmuir-Blodgett lithog. and subsequent radical polymerization) REFERENCE COUNT: THERE ARE 35 CITED REFERENCES AVAILABLE FOR 35 THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 2 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2007:221012 HCAPLUS Full-text DOCUMENT NUMBER: 146:442107

TITLE: Highly controlled living radical polymerization

through dual activation of

organobismuthines

AUTHOR(S): Yamago, Shigeru; Kayahara, Eiichi; Kotani,

Masashi; Ray, Biswajit; Kwak, Yungwan; Goto,

Atsushi; Fukuda, Takeshi

CORPORATE SOURCE: Div. Mol. Mater. Sci., Grad. Sch. Sci., Osaka City

Univ., Osaka, 558-8585, Japan

SOURCE: Angewandte Chemie, International Edition (2007),

46(8), 1304-1306

CODEN: ACIEF5; ISSN: 1433-7851

PUBLISHER: Wiley-VCH Verlag GmbH & Co. KGaA

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 146:442107

ED Entered STN: 01 Mar 2007

AB Organobismuthines promote highly controlled living radical polymerization through two activation mechanisms, namely, thermal generation and degenerative transfer. Both conjugated and nonconjugated vinyl monomers are polymerized to give well-defined polymers with predetd. mol. weight (Mn) and low polydispersity index (PDI).

IT 115191-52-1P

(highly controlled living radical polymerization of vinyl monomers through dual activation of organobismuthines)

RN 115191-52-1 HCAPLUS

CN Propanoic acid, 2-methyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]-, methyl ester (CA INDEX NAME)

CC 35-3 (Chemistry of Synthetic High Polymers)

 $\operatorname{ST}$   $\,$  vinyl monomer living radical polymn organobismuthine activation mechanism

IT Polymerization

Polymerization catalysts Polymerization kinetics

(living, radical; highly controlled living radical polymerization of vinyl monomers through dual activation of organobismuthines)

IT 934733-52-5P 934733-53-6P

(highly controlled living radical polymerization of vinyl monomers through dual activation of organobismuthines)

IT 100-42-5, Styrene, reactions

(highly controlled living radical polymerization of vinyl monomers through dual activation of organobismuthines)

IT 6180-99-0DP, Tributyltin deuteride, reaction products with polystyrene and organobismuthines 9003-39-8P, Poly(N-vinylpyrrolidone) 9003-49-0P, Poly(butyl acrylate) 9003-53-6DP, Polystyrene, reaction products with organobismuthines and tributyltin deuteride 9003-53-6P, Polystyrene 9011-14-7P, Poly(methyl methacrylate)

21735-64-8P 25189-55-3P, Poly(N-isopropylacrylamide) 115191-52-1P 733045-97-1P, Styrene-N-vinylpyrrolidone diblock copolymer 934733-52-5DP, reaction products with polystyrene and tributyltin deuteride

(highly controlled living radical polymerization of vinyl monomers through dual activation of organobismuthines)

IT 78-82-0, Isobutyronitrile 80-62-6 141-32-2 547-63-7, Methyl isobutyrate 2210-25-5 4540-16-3 7529-35-3, Dimethylbromobismuth 39248-62-9, Diphenylbismuth bromide

(preparation of organobismuthines for highly controlled living radical polymerization of vinyl monomers through dual activation of organobismuthines)

REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 3 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2006:1183825 HCAPLUS Full-text

DOCUMENT NUMBER: 146:101482

TITLE: Stabilization of polystyrene thin films against dewetting by silsesquioxane-terminated polystyrene

additives

AUTHOR(S): Miyamoto, Kyota; Hosaka, Nao; Otsuka, Hideyuki;

Takahara, Atsushi

CORPORATE SOURCE: Graduate School of Engineering, Kyushu University,

Higashi-ku, Fukuoka, 812-8581, Japan

SOURCE: Chemistry Letters (2006), 35(10), 1098-1099

CODEN: CMLTAG; ISSN: 0366-7022

PUBLISHER: Chemical Society of Japan

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 146:101482

ED Entered STN: 10 Nov 2006

AB A polyhedral oligomeric silsesquioxane (POSS)-containing initiator for nitroxide-mediated radical polymerization was synthesized to prepare organic-inorg. hybrid polymers (PS-POSS), which are polystyrene (PS) with a POSS end group. PS-POSS were well dispersed in PS thin films and provided thermal stability to films against dewetting.

IT 917594-78-6P

(stabilization of polystyrene thin films against dewetting by silsesquioxane-terminated polystyrene additives prepared by using polyhedral oligomeric silsesquioxane-containing nitroxide derivative as radical polymerization catalyst)

RN 917594-78-6 HCAPLUS

CN Carbamic acid, N-[3-[[(3,5,7,9,11,13,15-heptacyclopentylpentacyclo[9.5 .1.13,9.15,15.17,13]octasiloxan-1-yl)oxy]dimethylsilyl]propyl]-, 2-[(4-methoxy-2,2,6,6-tetramethyl-1-piperidinyl)oxy]-2-phenylethyl ester (CA INDEX NAME)

PAGE 1-A

PAGE 2-A

- CC 37-3 (Plastics Manufacture and Processing)
- ST polyhedral oligomeric silsesquioxane terminated polystyrene prepn; nitroxide deriv oligomeric silsesquioxane group initiator polystyrene polymn
- IT Wetting

(dewetting; stabilization of polystyrene thin films against dewetting by silsesquioxane-terminated polystyrene additives prepared by using polyhedral oligomeric silsesquioxane-containing nitroxide derivative as radical polymerization datalyst)

IT Polymerization

Polymerization catalysts

(radical; stabilization of polystyrene thin films against dewetting by silsesquioxane-terminated polystyrene additives prepared by using polyhedral oligomeric silsesquioxane-containing nitroxide derivative as radical polymerization catalyst)

IT 917594-78-6P

(stabilization of polystyrene thin films against dewetting by silsesquioxane-terminated polystyrene additives prepared by using polyhedral oligomeric silsesquioxane-containing nitroxide derivative as radical polymerization catalyst)

IT 9003-53-6DP, Polystyrene, polyhedral oligomeric silsesquioxane group-terminated

(stabilization of polystyrene thin films against dewetting by silsesquioxane-terminated polystyrene additives prepared by using polyhedral oligomeric silsesquioxane-containing nitroxide derivative as radical polymerization catalyst)

IT 9003-53-6, Polystyrene

(stabilization of polystyrene thin films against dewetting by silsesquioxane-terminated polystyrene additives prepared by using polyhedral oligomeric silsesquioxane-containing nitroxide derivative as radical polymerization catalyst)

IT 352538-81-9 491588-90-0

(stabilization of polystyrene thin films against dewetting by silsesquioxane-terminated polystyrene additives prepared by using polyhedral oligomeric silsesquioxane-containing nitroxide derivative as radical polymerization catalyst)

REFERENCE COUNT: 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 4 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2006:681741 HCAPLUS Full-text

DOCUMENT NUMBER: 145:293386

TITLE: Synthesis of poly(methyl methacrylate)-b-

polystyrene containing a crown ether unit at the junction point via combination of atom transfer radical polymerization and nitroxide mediated

radical polymerization routes

AUTHOR(S): Altintas, Ozcan; Yilmaz, Ismail; Hizal, Gurkan;

Tunca, Umit

CORPORATE SOURCE: Department of Chemistry, Istanbul Technical

University, Istanbul, 34469, Turk.

SOURCE: Journal of Polymer Science, Part A: Polymer

Chemistry (2006), 44(10), 3242-3249

CODEN: JPACEC; ISSN: 0887-624X

PUBLISHER: John Wiley & Sons, Inc.

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 14 Jul 2006

Poly(Me methacrylate)-b-polystyrene (PMMA-b-PS) containing a benzo-15-crown-5 unit at the junction point was prepared by combining atom transfer radical polymerization and nitroxide-mediated radical polymerization For this purpose, 6,7,9,10,12,13,15,16-octahydro-5,8,11,14,17-pentaoxa- benzocyclopentadecene-2-carboxylic acid 3-(2-bromo-2-methyl- propionyloxy)-2-methyl-2-[2-phenyl-2-(2,2,6,6-tetramethyl-piperidin-1- yloxy)-ethoxycarbonyl]-Pr ester (I) was synthesized and used as an initiator in atom transfer radical polymerization of Me methacrylate in the presence of CuCl and pentamethyldiethylenetriamine at 60°. A linear behavior was observed in both plots of In([M]0/[M]) vs. time and Mn,GPC vs. conversion indicating that the polymerization proceeded in a controlled/living manner. Thus obtained PMMA precursor was used as a macroinitiator in nitroxide-mediated radical polymerization of styrene (St) at 125° to give well-defined PMMA-b-PS with crown ether per chain. Kinetic data were also obtained for copolymn. Moreover, potassium picrate (K+ picrate) complexation of I and PMMA-b-PS copolymer was studied.

IT 908128-11-0P

(initiator; synthesis of poly(Me methacrylate)-b-

polystyrene containing crown ether unit at junction point via combination of atom transfer radical polymerization and nitroxide mediated radical polymerization routes)

RN 908128-11-0 HCAPLUS

CN 1,4,7,10,13-Benzopentaoxacyclopentadecin-15-carboxylic acid, 2,3,5,6,8,9,11,12-octahydro-, 2-[(2-bromo-2-methyl-1-oxopropoxy)methyl]-2-methyl-3-oxo-3-[2-phenyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethoxy]propyl ester (CA INDEX NAME)

CC 35-3 (Chemistry of Synthetic High Polymers)

ST polymethyl methacrylate polystyrene diblock copolymer crown ether nitroxide initiator; atom transfer radical polymn polymethyl methacrylate polystyrene diblock copolymer

IT Polymerization

Polymerization catalysts

Polymerization kinetics

(atom transfer, radical; synthesis of poly(Me methacrylate)-b-polystyrene containing crown ether unit at junction point via combination of atom transfer radical polymerization and nitroxide mediated radical polymerization routes)

IT 908128-11-0P

(initiator; synthesis of poly(Me methacrylate)-b-polystyrene containing crown ether unit at junction point via combination of atom transfer radical polymerization and nitroxide mediated radical polymerization routes)

9011-14-7DP, Poly(methyl methacrylate), reaction products with nitroxide-containing benzo-15-crown-5 derivs. 908128-11-0DP, reaction products with poly(Me methacrylate)

(macroinitiator; synthesis of poly(Me methacrylate)-b-polystyrene containing crown ether unit at junction point via combination of atom transfer radical polymerization and nitroxide mediated radical

polymerization

routes)

REFERENCE COUNT: 24 THERE ARE 24 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L68 ANSWER 5 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2006:632708 HCAPLUS Full-text

DOCUMENT NUMBER: 145:231158

TITLE: Exfoliated Block Copolymer/Silicate Nanocomposites

by One-Pot, One-Step in-Situ Living Polymerization

from Silicate-Anchored Multifunctional

Initiator

AUTHOR(S): Di, Jianbo; Sogah, Dotsevi Y.

CORPORATE SOURCE: Baker Laboratory, Department of Chemistry and

Chemical Biology, Cornell University, Ithaca, NY,

14853-1301, USA

SOURCE: Macromolecules (2006), 39(15), 5052-5057

CODEN: MAMOBX; ISSN: 0024-9297

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 30 Jun 2006

AB Poly(styrene-b-caprolactone)/silicate nanocomposites were prepared via one-pot, one-step in-situ living polymerization from a silicate-anchored bifunctional initiator. The random dispersion of the silicate layers in the polymer matrix was confirmed by both XRD and STEM. The polymer chains were attached to the surface of the silicate layers at the junction between the two blocks. SEC and NMR confirmed the block structure of the polymer. Through simultaneous incorporation of the initiator and benzyltrimethylammonium salt as a noninitiator into the silicate nanocomposites containing higher mol. weight polymers were obtained. The mol. wts. of the polymers and the silicate content of the nanocomposites were also controlled. Characterization by XRD and DSC showed that the poly(caprolactone) segment existed in a crystalline state.

IT 887369-62-2P

(ATRP initiator, clay anchored; preparation of multifunctional initiator for living polymerization to prepare exfoliated block copolymer/silicate nanocomposites)

RN 887369-62-2 HCAPLUS

CN Benzenemethanaminium, 4-[2-hydroxy-1-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl]-N,N,N-trimethyl-, chloride (9CI) (CA INDEX NAME)

● c1 -

CC 37-6 (Plastics Manufacture and Processing)
 Section cross-reference(s): 35

ST styrene caprolactone block copolymer silicate nanocomposite multifunctional initiator ATRP

IT Polymerization

(atom transfer, living, radical; exfoliated block copolymer/silicate nanocomposites by one-pot, one-step in-situ living polymerization from silicate-anchored multifunctional initiator)

IT Intercalation

Nanocomposites

(exfoliated block copolymer/silicate nanocomposites by one-pot, one-step in-situ living polymerization from silicate-anchored multifunctional initiator)

IT Silicates, preparation

(intercalation product with  $\epsilon\text{-Caprolactone-styrene}$  diblock copolymer; exfoliated block copolymer/silicate nanocomposites by one-pot, one-step in-situ living polymerization from silicate-anchored

10/519,030 multifunctional initiator) ΙT 887369-62-2P (ATRP initiator, clay anchored; preparation of multifunctional initiator for living polymerization to prepare exfoliated block copolymer/silicate nanocomposites) ΙT 97-93-8, Triethyl aluminum, uses (exfoliated block copolymer/silicate nanocomposites by one-pot, one-step in-situ living polymerization from silicate-anchored multifunctional initiator) 1318-93-0DP, Montmorillonite, sodium-exchanged, intercalation product ΙT with  $\epsilon$ -Caprolactone-styrene diblock copolymer 725712-80-1DP, ε-Caprolactone-styrene diblock copolymer, intercalation product with sodium-exchanged montmorillonite (exfoliated block copolymer/silicate nanocomposites by one-pot, one-step in-situ living polymerization from silicate-anchored multifunctional initiator) 75-50-3, Trimethylamine, reactions 196930-68-4 ΙT (preparation of multifunctional initiator for living polymerization to prepare exfoliated block copolymer/silicate nanocomposites) 100-59-4, Phenyl magnesiumchloride ΤT (preparation of multifunctional initiator for living polymerization to prepare exfoliated block copolymer/silicate nanocomposites) REFERENCE COUNT: 38 THERE ARE 38 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L68 ANSWER 6 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2005:1108314 HCAPLUS Full-text DOCUMENT NUMBER: 144:433168 New and improved routes to alkoxyamine TITLE: initiators for controlled radical polymerisation AUTHOR(S): Thiessen, Wladimir; Wolff, Thomas CORPORATE SOURCE: Institut fuer Physikalische Chemie und Elektrochemie, Technische Universitaet Dresden, Dresden, D-01062, Germany SOURCE: Designed Monomers and Polymers (2005), 8(5), 397-407 CODEN: DMPOF3; ISSN: 1385-772X PUBLISHER: VSP DOCUMENT TYPE: Journal LANGUAGE: English Entered STN: 17 Oct 2005 AΒ O-substituted, hydroxylamines were synthesized in almost quant. yields using ultrasound and Mn(OAc)3 (in substance or regenerated in situ by an oxidizing agent) as electron-transfer agent. Two novel synthetic methods for the synthesis of alkoxyamines from organic halides and stable nitroxide radicals are introduced. One of them utilizes cyancobalamine (vitamin B12), the other SmI2 as catalyst and magnesium metal as reductive agent. The yields are good for the first method and excellent for the second. The initiators were tested in controlled radical polymerization of styrene, Me methacrylate and 4vinylpyridine. ΙT 115191-52-1P (production of alkoxyamine initiators using

CN Propanoic acid, 2-methyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]-, methyl ester (CA INDEX NAME)

RN

115191-52-1 HCAPLUS

Mn(OAc)3 as electron-transfer agent for controlled radical polymerization)

```
CC
     35-3 (Chemistry of Synthetic High Polymers)
     Section cross-reference(s): 67
ST
     alkoxyamine initiator controlled radical polymn
ΙT
     Electron exchangers
     Oxidizing agents
     Sound and Ultrasound
        (production of alkoxyamine initiators for
        controlled radical polymerization)
ΙT
     Polymerization catalysts
        (radical; production of alkozyamine initiators for
        controlled radical polymerization)
ΙT
     993-02-2, Manganese triacetate
        (electron-transfer agent; production of alkoxyamine
        initiators using Mn(OAc)3 as electron-transfer agent for
        controlled radical polymerization)
ΙT
     9011-14-7P, Polymethylmethacrylate 25232-41-1P, Poly(4-
     vinylpyridine)
        (polymerization of Me methacrylate using alkoxyamine
        initiators as controlled radical polymerization catalyst)
     9003-53-6P, Polystyrene
ΤТ
        (polymerization of styrene using alkoxyamine initiators
        as controlled radical polymerization catalyst)
ΙT
     115191-52-1P 213699-59-3P
        (production of alkoxyamine initiators using
        Mn(OAc)3 as electron-transfer agent for controlled radical polymerization)
     80-62-6, Methylmethacrylate 100-42-5, Styrene, reactions 2564-83-2
ΙT
     2628-16-2
               188526-94-5, tert-Butyl 1-(diethoxyphosphinyl)-2,2-
     dimethylpropyl nitroxide
        (production of alkozyamine initiators using
        Mn(OAc)3 as electron-transfer agent for controlled radical polymerization)
     144-55-8, Sodium bicarbonate, reactions
ΤT
        (production of alkoxyamine initiators using
        Mn(OAc)3 as electron-transfer agent for controlled radical polymerization)
ΙT
     32248-43-4, Samarium diiodide
        (production of alkoxyamine initiators using SmI2 as
        catalyst for controlled radical polymerization)
ΙT
     224575-62-6P
        (production of alkoxyamine initiators using SmI2 as
        catalyst for controlled radical polymerization)
ΙT
     68-19-9, Vitamin B12
        (production of alkoxyamine initiators using as
        vitamin B12 catalyst for controlled radical polymerization)
ΙT
     103-63-9P
        (production of alkoxyamine initiators using as
        vitamin B12 catalyst for controlled radical polymerization)
     154554-67-3P, 2,2,6,6-Tetramethyl-1-(1-phenylethoxy)piperidine
ΤТ
        (production of alkoxyamine initiators using as
        vitamin B12 catalyst for controlled radical polymerization)
```

IT 12125-02-9, Ammonium chloride, reactions

(production of alkoxyamine initiators using as

vitamin B12 catalyst for controlled radical polymerization)

IT 7439-95-4, Magnesium, reactions

(reductive agent; production of alkoxyamine

initiators using SmI2 as catalyst for controlled radical

polymerization)

REFERENCE COUNT:

20 THERE ARE 20 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L68 ANSWER 7 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2005:609683 HCAPLUS Full-text

DOCUMENT NUMBER: 143:286754

TITLE: Nitroxide-Mediated Polymerization of

N-Isopropylacrylamide: Electrospray Ionization

Mass Spectrometry, Matrix-Assisted Laser Desorption Ionization Mass Spectrometry, and Multiple-Angle Laser Light Scattering Studies on Nitroxide-Terminated Poly-N-isopropylacrylamides

AUTHOR(S): Schulte, Tobias; Siegenthaler, Kai Oliver;

Luftmann, Heinrich; Letzel, Matthias; Studer,

Armido

CORPORATE SOURCE: Organisch-Chemisches Institut, Westfaelische

Wilhelms-Universitaet Muenster, Muenster, D-48149,

Germany

SOURCE: Macromolecules (2005), 38(16), 6833-6840

CODEN: MAMOBX; ISSN: 0024-9297

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 15 Jul 2005

Nitroxide-mediated controlled living free radical polymerization of N-isopropylacrylamide using highly sterically hindered 2,2,6,6-tetraethylpiperidin-4-on-N-oxyl 1 is described. In addition, an improved synthesis for nitroxide 1 is presented. Poly-N-isopropylacrylamides (PNIPAMs) prepared are analyzed by multiple-angle laser light scattering. Moreover, the nitroxide-terminated PNIPAMs are characterized using electrospray ionization mass spectrometry, matrix-assisted laser desorption ionization time-of-flight mass spectrometry (MALDI-TOF-MS), and Fourier transform ion cyclotron MALDI-MS. Careful MS anal. reveals that chain-end degradation of nitroxide-terminated PNIPAMs occurs during MALDI anal. A mechanism for chain end degradation is presented.

IT 686778-17-6P

(preparation of tetraethylpiperidinonoxyl and alkoxyamineinitiated living radical polymerization of isopropylacrylamide and study of structure and chain-end degradation of nitroxide-terminated polyisopropylacrylamide)

RN 686778-17-6 HCAPLUS

CN 4-Piperidinone, 2,2,6,6-tetraethyl-1-(1-phenylethoxy)- (CA INDEX NAME)

35-4 (Chemistry of Synthetic High Polymers) Section cross-reference(s): 27 ST living radical polymn isopropylacrylamide alkoxyamine initiator; nitroxide terminated living polyisopropylacrylamide mass spectrometry light scattering Polymer degradation ΙT (chain end degradation mechanism; preparation of tetraethylpiperidinonoxyl and alkoxyamine-initiated living radical polymerization of isopropylacrylamide and study of structure and chain-end degradation of nitroxide-terminated polyisopropylacrylamide) Polvmerization ΙT Polymerization catalysts (living, radical; preparation of tetraethylpiperidinonoxyl and alkoxyamine-initiated living radical polymerization of isopropylacrylamide and study of structure and chain-end degradation of nitroxide-terminated polyisopropylacrylamide) Electrospray ionization mass spectrometry ΤТ Light scattering Photoionization mass spectrometry (preparation of tetraethylpiperidinonoxyl and alkoxyamineinitiated living radical polymerization of isopropylacrylamide and study of structure and chain-end degradation of nitroxide-terminated polyisopropylacrylamide using) 686778-17-6P ΙT (preparation of tetraethylpiperidinonoxyl and alkoxyamineinitiated living radical polymerization of isopropylacrylamide and study of structure and chain-end degradation of nitroxide-terminated polvisopropvlacrylamide) ΙT 25189-55-3DP, N-Isopropylacrylamide homopolymer, alkoxyamineterminated 686778-08-5DP, reaction products with poly(N-isopropylacrylamide) (preparation of tetraethylpiperidinonoxyl and alkoxyamineinitiated living radical polymerization of isopropylacrylamide and study of structure and chain-end degradation of nitroxide-terminated polyisopropylacrylamide) ΙT 760-21-4, 2-Ethyl-1-butene 1189-71-5, Chlorosulfonyl isocyanate 24424-99-5, Di-tert-butyl dicarbonate (preparation of tetraethylpiperidinonoxyl and alkoxyamineinitiated living radical polymerization of isopropylacrylamide and study of structure and chain-end degradation of nitroxide-terminated polyisopropylacrylamide) ΤТ 89894-89-3P 864062-67-9P 864062-68-0P 864062-69-1P 864062-70-4P (preparation of tetraethylpiperidinonoxyl and alkoxyamineinitiated living radical polymerization of isopropylacrylamide and study of structure and chain-end degradation of nitroxide-terminated polyisopropylacrylamide) ΙT 686778-08-5P (preparation of tetraethylpiperidinonoxyl and alkoxyamineinitiated living radical polymerization of isopropylacrylamide and study of structure and chain-end degradation of nitroxide-terminated polyisopropylacrylamide) REFERENCE COUNT: 47 THERE ARE 47 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE

L68 ANSWER 8 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2005:364390 HCAPLUS Full-text

RE FORMAT

DOCUMENT NUMBER: 143:59393

TITLE: Tin-free radical alkoxyamine addition and

> isomerization reactions by using the persistent radical effect: Variation of the alkoxyamine

structure

AUTHOR(S): Molawi, Kian; Schulte, Tobias; Siegenthaler, Kai

> Oliver; Wetter, Christian; Studer, Armido Organisch Chemisches-Institut Westfaelische

CORPORATE SOURCE: Wilhelms-Universitaet, Muenster, 48149, Germany

Chemistry--A European Journal (2005), 11(8),

2335-2350

CODEN: CEUJED; ISSN: 0947-6539

Wiley-VCH Verlag GmbH & Co. KGaA PUBLISHER:

DOCUMENT TYPE: Journal English LANGUAGE:

OTHER SOURCE(S): CASREACT 143:59393

Entered STN: 28 Apr 2005

AΒ Various C-centered radicals can efficiently be generated through thermal C-Obond homolysis of alkoxyamines. This method is used to perform environmentally benign radical cyclization and intermol. addition reactions. These alkoxyamine isomerizations and intermol. carboaminoxylations are mediated by the persistent radical effect (PRE). In the paper, the effect of the variation of the alkoxyamine structure-in particular steric effects in the nitroxide moiety-on the outcome of the PRE mediated radical reactions will be discussed. Fourteen different nitroxides were used in the studies. It will be shown that reaction times can be shortened about 100 times upon careful tuning of the alkoxyamine structure. Activation energies for the C-O-bond homolysis of the various alkoxyamines are provided. The kinetic data are used to explain the reaction outcome of the PRE-mediated processes.

ΙT 270901-55-8P

SOURCE:

(tin-free radical alkoxyamine addition and isomerization reactions by using the persistent radical effect)

RN 270901-55-8 HCAPLUS

6-Heptenoic acid, 2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]-, CN 1,1-dimethylethyl ester (CA INDEX NAME)

CC 22-4 (Physical Organic Chemistry) TT 270901-55-8P 270901-57-0P 270901-58-1P 270901-83-2P 270901-84-3P 853886-27-8P 853886-33-6P 853886-36-9P 853886-37-0P 853886-40-5P 853886-44-9P 853886-48-3P 853886-51-8P 853886-55-2P 853886-56-3P 853886-57-4P 853886-58-5P 853886-59-6P **853886-60-9P** 853886-65-4P 853886-66-5P 853886-67-6P 853886-68-7P 853886-69-8P 853886-70-1P 853886-77-8P 853886-79-0P 853886-81-4P 853886-82-5P

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(tin-free radical alkoxyamine addition and isomerization reactions by
       using the persistent radical effect)
    853886-26-7P
        (tin-free radical alkoxyamine addition and isomerization reactions by
       using the persistent radical effect)
    270901-60-5P 270901-70-7P 270901-72-9P
ΙT
                                                270901-89-8P
    270901-90-1P 270901-93-4P 270901-95-6P 675879-82-0P
    853886-20-1P 853886-21-2P 853886-22-3P 853886-23-4P
    853886-24-5P 853886-25-6P 853886-28-9P 853886-29-0P
    853886-30-3P 853886-31-4P 853886-32-5P 853886-34-7P
    853886-35-8P 853886-38-1P 853886-39-2P 853886-41-6P
    853886-42-7P 853886-43-8P 853886-45-0P 853886-46-1P
    853886-47-2P 853886-49-4P 853886-50-7P 853886-52-9P
    853886-53-0P 853886-54-1P 853886-61-0P 853886-62-1P
    853886-63-2P 853886-64-3P 853886-71-2P 853886-72-3P
    853886-73-4P 853886-74-5P 853886-75-6P 853886-76-7P
    853886-78-9P 853886-80-3P 853886-83-6P 853886-84-7P
    853886-85-8P 853886-86-9P 853904-90-2P
        (tin-free radical alkoxyamine addition and isomerization reactions by
       using the persistent radical effect)
                              THERE ARE 48 CITED REFERENCES AVAILABLE FOR
REFERENCE COUNT:
                        48
                              THIS RECORD. ALL CITATIONS AVAILABLE IN THE
                              RE FORMAT
L68 ANSWER 9 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:
                       2004:686753 HCAPLUS Full-text
DOCUMENT NUMBER:
                        142:135004
TITLE:
                        A novel method for the synthesis of
                        alkoxyamine initiators for
                        nitroxide-mediated radical polymerization using
                        Mn(OAc)3 as electron-transfer reagent
                        Krause, T.; Habicher, W. D.; Messerschmidt, M.;
AUTHOR(S):
                        Voit, B. I.
CORPORATE SOURCE:
                        Institut fuer Organische Chemie, Technische
                        Universitaet Dresden, Dresden, 01062, Germany
                        Designed Monomers and Polymers (2004), 7(4),
SOURCE:
                        391-397
                        CODEN: DMPOF3; ISSN: 1385-772X
                        VSP BV
PUBLISHER:
DOCUMENT TYPE:
                        Journal
LANGUAGE:
                        English
    Entered STN: 23 Aug 2004
ED
     O-substituted hydroxylamines which are known to initiate and promote
AB
     nitroxide-mediated radical polymerization were synthesized in high yield by a
     novel and facile synthetic approach. Using the favorable Mn(OAc)3 as electron
     transfer agent in the in situ generation of benzylic radicals and their
     trapping by nitroxide radicals provides a new powerful and more economic way
     to the desired alkoxyamine initiators for this controlled, living radical
     polymerization TEMPO, TIPNO and the protected tris-hydroxy derivative of
     TIPNO were used to synthesize the appropriate alkoxyamines.
ΙT
    154554-67-3P
        (synthesis of alkoxyamine initiators for
       nitroxide-mediated radical polymerization using Mn(OAc)3 as
       electron-transfer reagent)
RN
    154554-67-3 HCAPLUS
CN
    Piperidine, 2,2,6,6-tetramethyl-1-(1-phenylethoxy)- (CA INDEX NAME)
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CC 35-3 (Chemistry of Synthetic High Polymers)

IT Polymerization catalysts

(synthesis of alkoxyamine initiators for

nitroxide-mediated radical polymerization using Mn(OAc)3 as electron-transfer reagent)

IT 993-02-2, Manganese triacetate

(synthesis of alkoxyamine initiators for

nitroxide-mediated radical polymerization using Mn(OAc)3 as electron-transfer reagent)

IT 154554-67-3P 330938-14-2P 824430-24-2P 824430-25-3P (synthesis of alkoxyamine initiators for

nitroxide-mediated radical polymerization using Mn(OAc)3 as electron-transfer reagent)

IT 100-42-5, reactions 2564-83-2 2628-16-2 61015-94-9 95418-58-9 270901-77-4

(synthesis of alkoxyamine initiators for

nitroxide-mediated radical polymerization using Mn(OAc)3 as
electron-transfer reagent)

REFERENCE COUNT: 17 THERE ARE 17 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE REFORMAT

L68 ANSWER 10 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2003:1007898 HCAPLUS <u>Full-text</u>

DOCUMENT NUMBER: 140:60153

TITLE: One-pot process for preparation of hydroxy-functionalized alkoxyamine

initiator and its use

INVENTOR(S): Detrembleur, Christophe; Gross, Thomas; Meyer,

Rolf-Volker

PATENT ASSIGNEE(S): Bayer Aktiengesellschaft, Germany SOURCE: U.S. Pat. Appl. Publ., 12 pp.

CODEN: USXXCO

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND DATE		APPLICATION NO.	DATE		
US 2003236368	A1	20031225	US 2003-464927	20030619		
US 6686424	B2	20040203				
EP 1375457	A1	20040102	EP 2002-13949	20020625		
R: AT, BE, CH,	DE, DK,	, ES, FR, GB	, GR, IT, LI, LU, NL,	SE, MC,		
PT, IE, SI,	LT, LV,	, FI, RO, MK	, CY, AL, TR			
CA 2432964	A1	20031225	CA 2003-2432964	20030620		
JP 2004107320	A	20040408	JP 2003-181155	20030625		
PRIORITY APPLN. INFO.:			EP 2002-13949	A 20020625		

OTHER SOURCE(S): MARPAT 140:60153

ED Entered STN: 28 Dec 2003

Functional alkoxyamines of the general formula HOC(R1)HCR2R3ONR4R5 [R1-3 = H, C1-20 (cyclo)alkyl, C6-24 aryl, halogen, cyano, C1-20 (cyclo)alkyl ester, C1-20 (cyclo) alkylamido, C6-24 aryl ester, C6-24 arylamido group; R4-5 = optionally substituted C1-18 alk(e)nyl or alkynyl, C3-12 cycloalkyl, heterocycloalkyl, C6-24 aryl, etc., R4 and R5 together can make a ring, and optionally with N, O or S] are prepared by: (1) reacting an oxidizing agent with a sterically hindered secondary amine to produce an aqueous phase and a nitroxyl radical, and (2) removing the aqueous phase and adding to the nitroxyl radical one or more vinyl monomer(s) conforming to a formula and a system which produces free radicals. Also disclosed is a controlled radical process for polymerizing monomers using the functional alkoxyamine. Thus, a typical alkoxyamine such as 1-phenyl-1-(2',2',6',6'-tetramethyl-1'-piperidinyloxy)-2-hydroxyethane could be prepared from the reaction of Oxone with 2,2,6,6-tetramethylpiperidine, then with styrene and H2O2.

IT 161776-41-6P

(one-pot process for preparation of functionalized alkoxyamine initiator and its use)

RN 161776-41-6 HCAPLUS

CN Benzeneethanol,  $\beta$ -[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]- (CA INDEX NAME)

IC ICM C08F002-00

CS C08F004-00; C07D211-20; C07D223-08; C07D207-46; C07C255-62; C07C239-12

CC 35-3 (Chemistry of Synthetic High Polymers)

IT Polymerization catalysts

(radical; one-pot process for preparation of functionalized alkoxyamine initiator and its use)

IT 161776-41-6P 637334-58-8P

(one-pot process for preparation of functionalized alkoxyamine initiator and its use)

IT 9003-53-6P, Polystyrene 9003-54-7P, Acrylonitrile-styrene copolymer 126639-07-4P, Acrylonitrile-methyl methacrylate-styrene block copolymer

(one-pot process for preparation of functionalized alkoxyamine initiator and its use)

IT 2564-83-2P, 2,2,6,6-Tetramethylpiperidine-N-oxide (one-pot process for preparation of functionalized alkoxyamine initiator and its use)

IT 100-42-5, Styrene, reactions 768-66-1, 2,2,6,6-Tetramethylpiperidine 826-36-8, 2,2,6,6-Tetramethyl-4-piperidone (one-pot process for preparation of functionalized alkoxyamine

initiator and its use)

IT 7722-84-1, Hydrogen peroxide, reactions

(one-pot process for preparation of functionalized alkoxyamine initiator and its use)

alkoxyamine initiator and its use)

L68 ANSWER 11 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2003:961305 HCAPLUS Full-text

DOCUMENT NUMBER: 140:146562

TITLE: Hydroxy- and Silyloxy-Substituted TEMPO

Derivatives for the Living Free-Radical

Polymerization of Styrene and n-Butyl Acrylate: Synthesis, Kinetics, and Mechanistic Studies Knoop, Christoph Alexander; Studer, Armido

AUTHOR(S): Knoop, Christoph Alexander; Studer, Armido CORPORATE SOURCE: Department of Chemistry, Philipps-University

Marburg, Marburg, 35032, Germany

SOURCE: Journal of the American Chemical Society (2003),

125(52), 16327-16333

CODEN: JACSAT; ISSN: 0002-7863

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 10 Dec 2003

The synthesis of new 2,2,6,6-tetramethylpiperidin-1-oxyl (TEMPO) styryl AB derivs. as mediators for the living free-radical polymerization is described. Two of the  $\alpha\text{-Me}$  groups at the 2- and 6-position of the parent TEMPO styryl alkoxyamine have been replaced by hydroxymethyl and silyloxymethyl groups. further increase the steric hindrance around the alkoxyamine oxygen atom, the remaining two Me groups have been substituted with larger Et groups. Styrene polymns. using hydroxy-substituted TEMPO derivs. are fast, but are not wellcontrolled. As previously shown for other OH-substituted alkoxyamines, intramol. H-bonding leads to an acceleration of the C-O bond homolysis and, hence, to an acceleration of the polymerization process. However, the OH groups also increase the alkoxyamine decomposition rate constant. The kinetics of the C-O bond homolysis have been determined using EPR spectroscopy. Decomposition studies have been conducted with the aid of 1H NMR spectroscopy. In contrast to the OH-substituted alkoxyamines, highly hindered silyloxysubstituted TEMPO alkoxyamines turned out to be excellent mediator/initiators for the controlled styrene polymerization Polystyrene with Mn of up to 80,000 g/mol and narrow polydispersities (PDI) has been prepared using the new alkoxyamines. Reactions have been conducted at 105°; however, even at 90° controlled but slow polymns. can be achieved. Furthermore, and more importantly, poly(Bu acrylates) with narrow PDIs (<1.15) have been prepared at 105° with the new alkoxyamines. Controlled acrylate polymerization can be conducted at temps. as low as 90°. The silylated alkoxyamines presented belong to the most efficient initiator/mediators for the controlled acrylate polymerization known to date. The effect of the addition of free nitroxide on the acrylate polymerization is discussed. Moreover, the synthesis of diblock copolymers with narrow PDIs is described.

IT 651351-97-2P

(synthesis, kinetics, and mechanistic studies of hydroxy- and silyloxy-substituted tetramethylpiperidinoxyl derivs. for living free-radical polymerization of styrene and n-Bu acrylate)

RN 651351-97-2 HCAPLUS

CN 2,6-Piperidinedicarboxylic acid, 2,6-dimethyl-1-(1-phenylethoxy)-, dimethyl ester, (2R,6R)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.

CC 35-3 (Chemistry of Synthetic High Polymers)

ST hydroxy silyloxy substituted tetramethylpiperidinoxyl deriv styrene living radical polymn; butyl acrylate living radical polymn hydroxy silyloxy tetramethylpiperidinoxyl deriv; kinetics living radical polymn hydroxy silyloxy tetramethylpiperidinoxyl deriv initiator

IT Polymerization catalysts

(living, radical; synthesis, kinetics, and mechanistic studies of hydroxy- and silyloxy-substituted tetramethylpiperidinoxyl derivs. for living free-radical polymerization of styrene and n-Bu acrylate)

IT 651351-97-2P 651351-99-4P 651352-01-1P 651352-03-3P 651352-05-5P 651352-07-7P

651352-09-9P 651352-11-3P

(synthesis, kinetics, and mechanistic studies of hydroxy- and silyloxy-substituted tetramethylpiperidinoxyl derivs. for living free-radical polymerization of styrene and n-Bu acrylate)

IT 651352-12-4P 651352-13-5P 651352-15-7P 651352-17-9P 651352-19-1P 651352-21-5P

651352-25-9P

(synthesis, kinetics, and mechanistic studies of hydroxy- and silyloxy-substituted tetramethylpiperidinoxyl derivs. for living free-radical polymerization of styrene and n-Bu acrylate)

REFERENCE COUNT:

THERE ARE 52 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 12 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2003:719523 HCAPLUS Full-text

52

DOCUMENT NUMBER: 139:246324

TITLE: Open-chain alkoxyamines and their corresponding

nitroxides for controlled low-temperature radical

polymerization

INVENTOR(S): Hintermann, Tobias; Nesvadba, Peter; Kramer,

Andreas; Fink, Jochen

PATENT ASSIGNEE(S): Ciba Specialty Chemicals Holding Inc., Switz.

SOURCE: PCT Int. Appl., 59 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003074572	A1	20030912	WO 2003-EP1895	20030225

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W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH,
             CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD,
             GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ,
             LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,
             NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ,
             TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,
             BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK,
             EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI,
             SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE,
             SN, TD, TG
     CA 2477728
                          Α1
                                 20030912
                                            CA 2003-2477728
                                                                     20030225
                                             AU 2003-212272
     AU 2003212272
                          Α1
                                 20030916
                                                                    20030225
     EP 1481012
                                20041201
                                            EP 2003-708135
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                                 20060517
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC,
             PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
     JP 2005519121
                                20050630
                                            JP 2003-573037
                          Τ
                                                                    20030225
     CN 1639201
                          Α
                                20050713
                                            CN 2003-805021
                                                                    20030225
     AT 326487
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                                20060615
                                            AT 2003-708135
                                                                    20030225
     ES 2263957
                          ТЗ
                                20061216
                                            ES 2003-708135
                                                                    20030225
     MX 2004PA08321
                                20041126
                                            MX 2004-PA8321
                                                                    20040827
     US 2005124814
                                20050609
                                            US 2004-506700
                                                                    20040902
                          Α1
     US 7297819
                          В2
                                20071120
PRIORITY APPLN. INFO.:
                                            EP 2002-405168
                                                                 A 20020305
                                            WO 2003-EP1895 W 20030225
```

OTHER SOURCE(S): MARPAT 139:246324

ED Entered STN: 14 Sep 2003

GΙ

AB Alkoxyamines I, II, and III [Y = O or NR; R = H or C1-18 alkyl; R7 and(or) R8 with R and N to which they are bonded form 5-6-membered ring; R1-3 = organic; R4 = C2-12 alkyl; R5, R6 = H, C1-18 alkyl, C2-18 alkenyl, benzyl, C5-12 cycloalkyl, or Ph; R7, R8 = H or organic; R9-14 = H, OH, SH, or organic; X = organic] are useful for enhancing the polymerization rates and monomer-to-polymer conversions of ethylenically unsatd. compds. at ≤100°. The

intermediate N-oxyl derivs., a composition of the N-oxyl derivs. with ethylenically unsatd. monomers and a free radical initiator  $X \bullet$ , as well as a process and their use for polymerization are also subjects of the present invention. I (R1, R2 = Me, R3, R4 = Et, R5-8 = H, X = PhCHMe) was manufactured by adding 13.9 g 3,3-diethyl-5,5-dimethylmorpholin-2-one N-oxyl to THF containing LiAlH4 at 0-10° and heating 5 h at reflux.

IT 332380-44-6P, 3,3,5,5-Tetramethyl-4-(1-phenylethoxy)morpholin-2-one

(initiator model compound precursor; open-chain hindered alkoxyamines and their corresponding nitroxides for controlled low-temperature radical polymerization)

RN 332380-44-6 HCAPLUS

CN 2-Morpholinone, 3,3,5,5-tetramethyl-4-(1-phenylethoxy)- (CA INDEX NAME)

IC ICM C08F004-00

ICS C07C239-20; C07D265-32; C07D265-34

CC 35-3 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 23, 25

ST open chain hindered alkoxyamine initiator unsatd monomer radical polymn; ethylhydroxydimethylethyl phenylethoxyaminobutanol manuf initiator unsatd monomer radical polymn

IT 332380-44-6P, 3,3,5,5-Tetramethyl-4-(1-phenylethoxy)morpholin-

2-one

(initiator model compound precursor; open-chain hindered alkoxyamines and their corresponding nitroxides for controlled low-temperature radical polymerization)

IT 213924-52-8P, Sodium 2-(2-hydroxy-1,1-dimethylethylamino)-2-phenylpropionate 264279-70-1P, 3,3-Diethyl-5,5-dimethyl-4-(1-phenylethoxy)morpholin-2-one 597555-36-7P,

3,3-Diisopropyl-5,5-dimethyl-4-(1-phenylethoxy)morpholin-2-one

597555-40-3P, 3-Pentyl-3,5,5-trimethyl-4-(1-

phenylethoxy) morpholin-2-one 597555-44-7P, 2,2-Dimethyl-1-(1-mathematical phenylethoxy)

phenylethoxy)-1-aza-4-oxaspiro[5.5]undecan-5-one 597555-45-8P,

2,2-Dimethyl-1-(1-phenylethoxy)-1-aza-4-oxaspiro[5.6]dodecan-5-one 597555-47-0P, 3-Phenyl-3,5,5-trimethyl-4-(1-

phenylethoxy)morpholin-2-one 597555-49-2P,

5,5-Dimethyl-3-ethyl-3-phenyl-4-(1-phenylethoxy)morpholin-2-one 597555-52-7P, 3-(4-Chlorophenyl)-3,5,5-trimethyl-4-(1-

phenylethoxy)morpholin-2-one

(precursor; open-chain hindered alkoxyamines and their corresponding nitroxides for controlled low-temperature radical

polymerization)

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L68 ANSWER 13 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2003:434535 HCAPLUS Full-text

DOCUMENT NUMBER: 139:22821

TITLE: Process for the synthesis of hindered amine ethers

from secondary amino oxides
Frey, Markus; Rast, Valerie

PATENT ASSIGNEE(S): Ciba Specialty Chemicals Holding Inc., Switz.

SOURCE: PCT Int. Appl., 56 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

INVENTOR(S):

	PATENT NO.								APPLICATION NO.  WO 2002-EP12957									
WO	2003	0459	19		A3		2004	0429										
	W:	ΑE,	AG,	AL,	AM,	ΑT,	ΑU,	AZ,	BA,	BB	, BG,	BR,	BY,	BZ,	CA,	CH,		
		CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ	, EC,	EE,	ES,	FI,	GB,	GD,		
		GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS	, JP,	ΚE,	KG,	KP,	KR,	KΖ,		
		LC,	LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD	, MG,	MK,	MN,	MW,	MX,	MZ,		
		NO,	NZ,	OM,	PH,	PL,	PT,	RO,	RU,	SC	, SD,	SE,	SG,	SI,	SK,	SL,		
		ΤJ,	TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US	, UZ,	VC,	VN,	YU,	ZA,	ZM,	ZW	
	RW:	GH,	GM,	ΚE,	LS,	MW,	MZ,	SD,	SL,	SZ	, TZ,	UG,	ZM,	ZW,	AM,	ΑZ,		
		BY,	KG,	KΖ,	MD,	RU,	ТJ,	TM,	ΑT,	BE	, BG,	CH,	CY,	CZ,	DE,	DK,		
		EE,	ES,	FI,	FR,	GB,	GR,	ΙE,	IT,	LU	, MC,	NL,	PT,	SE,	SK,	TR,		
		BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ	, GW,	ML,	MR,	ΝE,	SN,	TD,	ΤG	
CA	2464	107			A1		2003	0605		CA :	2002-	2464	107		2	0021	119	
AU	2002	3520.	57		A1 20030610				AU :	2002-	3520	57		2	0021	119		
EP	1463	717			A2 20041006				EP :	2002-	7877	31		2	0021	119		
	R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR	, IT,	LI,	LU,	NL,	SE,	MC,		
		PT,	ΙE,	SI,	LT,	LV,	FI,	RO,	MK,	CY	, AL,	TR,	BG,	CZ,	EE,	SK		
BR	2002	0144	29		Α		2004	1103	BR 2002-14429						20021119			
CN	1592	740			Α		2005	0309	CN 2002-823421 JP 2003-547371									
JP	2005	5169	05		Τ		2005	0609		JP :	2003-	5473	71		2	0021	119	
MX	2004	PA04	694		Α		2004	0819		MX :	2004-	PA46	94		2	0040	518	
US	2005	1040	42		A1		2005	0519		US :	2004-	4967	73		2	0040	524	
PRIORIT:	Y APP	LN.	INFO	. :						EP :	2001-	8111	43		A 2	0011	126	
										WO :	2002-	EP12	957	1	W 2	0021	119	

OTHER SOURCE(S): MARPAT 139:22821

ED Entered STN: 06 Jun 2003

AB Amine ethers of sterically hindered amines are obtained in good yield from the corresponding N-oxyl hindered amine precursor by reaction with a hydrocarbon in the presence of an organic hydroperoxide and an iodide. The products of present process find utility as polymerization regulators and/or light stabilizers for organic material. Thus, adding tert-Bu hydroperoxide (I; 70% aqueous solution) 6.2 to a stirred mixture of 2,2,6,6-tetramethylpiperidine-N-oxide (TEMPO) 5, ethylbenzene 34 and tetrabutylammonium iodide 0.12 g within 30 min, heating at 60° for 25 min until all of the TEMPO has reacted, cooling to 25°, stirring with a 10% aqueous solution of Na2SO3 until the disappearance of excess I, separating the aqueous phase, washing and drying over MgSO4 gave 1-(1-phenylethoxy)-2,2,6,6-tetramethylpiperidine.

IT 122586-81-6P

(process for synthesis of hindered amine ethers from secondary amino oxides)

RN 122586-81-6 HCAPLUS

CN 4-Piperidinone, 2,2,6,6-tetramethyl-1-(1-phenylethoxy)- (CA INDEX NAME)

TC ICM C07D211-94 ICS C07D401-14; C07D493-10; C07C239-20; C09K015-20; C08K005-3435; C08K005-3492; C08F002-38 CC 37-2 (Plastics Manufacture and Processing) ΤТ Oxidation catalysts (onium salts; process for synthesis of hindered amine ethers from secondary amino oxides) ΙT 117174-66-0P 122586-81-6P 122587-12-6P 154554-67-3P, 1-(1-Phenylethoxy)-2,2,6,6-tetramethylpiperidine243972-05-6P 244021-01-0P 264224-73-9P 335201-37-1P 378245-16-0P 378245-17-1P 378245-30-8P 434898-80-3P 437744-23-5P 437748-41-9P 538343-67-8P 538343-74-7P 538343-76-9P 538343-80-5P (process for synthesis of hindered amine ethers from secondary amino oxides)

L68 ANSWER 14 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2003:397228 HCAPLUS Full-text

DOCUMENT NUMBER: 139:117736

TITLE: Synthesis of reactive poly(vinyl oxazolones) via

nitroxide-mediated "living" free radical

polymerization

AUTHOR(S): Tully, David C.; Roberts, Michael J.;

Geierstanger, Bernhard H.; Grubbs, Robert B.

CORPORATE SOURCE: Genomics Institute, Novartis Research Foundation,

San Diego, CA, 92121, USA

SOURCE: Macromolecules (2003), 36(12), 4302-4308

CODEN: MAMOBX; ISSN: 0024-9297

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 25 May 2003

Low-polydispersity poly(vinyl oxazolone) has been prepared using nitroxide-mediated living free radical polymerization. Bulk homopolymn. of 2-vinyl-4,4-dimethyl-5-oxazolone (VDMO) in the presence of the  $\alpha$ -hydrido alkoxyamine initiator, N-(1,1-dimethylethyl)- $\alpha$ -(1-methylethyl)-N-(1-phenylethoxy)-benzenemethanamine, and the corresponding nitroxide proceeds to high conversion with polydispersities of less than 1.10. Accurate mol. weight control and low polydispersities (.apprx.1.04-1.10) were obtained on statistical copolymn. with styrene. Well-defined reactive statistical copolymers with polydispersities ranging from 1.05 to 1.30 were also prepared by copolymn. of oxazolone-functional monomers with acrylates, acrylamides, and N-vinylamides. Reactive block copolymers were prepared by polymerization of VDMO from poly(Bu acrylate) starting blocks as well as by polymerization of styrene from poly(VDMO) starting blocks. New polymers were prepared from

VDMO-containing polymers and copolymers by nucleophilic ring-opening of the pendant oxazolone rings with amines.

IT 154554-67-3

(synthesis of reactive poly(vinyl oxazolones) via nitroxide-mediated "living" free radical polymerization)

RN 154554-67-3 HCAPLUS

CN Piperidine, 2,2,6,6-tetramethyl-1-(1-phenylethoxy)- (CA INDEX NAME)

CC 35-4 (Chemistry of Synthetic High Polymers)

IT 2564-83-2, TEMPO 61015-94-9 154554-67-3 227000-59-1

(synthesis of reactive poly(vinyl oxazolones) via

nitroxide-mediated "living" free radical polymerization)

REFERENCE COUNT: 23 THERE ARE 23 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L68 ANSWER 15 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2003:282982 HCAPLUS Full-text

DOCUMENT NUMBER: 139:22546

TITLE: New Seven- and Eight-Membered Cyclic Alkoxyamines

for the Living Free Radical Polymerization

AUTHOR(S): Schulte, Tobias; Studer, Armido

CORPORATE SOURCE: Fachbereich Chemie der Universitaet Marburg,

Marburg, D-35032, Germany

SOURCE: Macromolecules (2003), 36(9), 3078-3084

CODEN: MAMOBX; ISSN: 0024-9297

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 13 Apr 2003

AB A straightforward synthesis of new seven- and eight-membered cyclic alkoxyamines from the corresponding lower homologous keto-alkoxyamines via ring-enlargement using TMS-diazomethane is described. The use of these ring-enlarged cyclic alkoxyamines as regulators/initiators for the radical polymerization of styrene and Bu acrylate is presented. Efficient controlled and living styrene polymerization (mol. weight of up to 40 000) can be obtained using the seven- and eight-membered alkoxyamine initiators. The influence of the ring-enlargement on the quality of the polymerization process (polymerization time, livingness, PDI) is discussed. The rate constant of the C-O bond cleavage of these new alkoxyamines was measured. In addition, the thermal decomposition of the alkoxyamines was studied. Furthermore, EPR data of the corresponding new nitroxides are presented.

IT 122586-81-6P

(new seven- and eight-membered cyclic alkoxyamines catalysts for living free radical polymerization)

RN 122586-81-6 HCAPLUS

CN 4-Piperidinone, 2,2,6,6-tetramethyl-1-(1-phenylethoxy)- (CA INDEX NAME)

```
CC
     35-3 (Chemistry of Synthetic High Polymers)
    prepn seven eight membered cyclic alkoxyamine radical polymn
ST
     catalyst; alkoxyamine living free radical polymn
     catalyst prepn
ΤТ
    Activation energy
        (homolysis; new seven- and eight-membered cyclic alkoxyamines
        catalysts for living free radical polymerization)
ΙT
     Polymerization
     Polymerization catalysts
        (living, radical; new seven- and eight-membered cyclic alkoxyamines
        catalysts for living free radical polymerization)
ΤТ
     Hyperfine coupling
     Reduction
     Ring enlargement
     Ring enlargement catalysts
        (new seven- and eight-membered cyclic alkoxyamines
        catalysts for living free radical polymerization)
     Decomposition kinetics
ΤТ
        (of catalysts; new seven- and eight-membered cyclic
        alkoxyamines catalysts for living free radical polymerization)
ΙT
     122586-81-6P
                   538376-93-1P
                                   538376-94-2P
                                                  538376-95-3P
        (new seven- and eight-membered cyclic alkoxyamines
        catalysts for living free radical polymerization)
ΙT
     132416-36-5P
                   538376-96-4P
                                  538376-97-5P
                                                  538376-98-6P
        (new seven- and eight-membered cyclic alkoxyamines
        catalysts for living free radical polymerization)
     2226-96-2P 2896-70-0P 538376-99-7P 538377-00-3P
ΙT
                                                             538377-01-4P
     538377-02-5P
                  538377-03-6P
                                   538377-04-7P
        (new seven- and eight-membered cyclic alkoxyamines
        catalysts for living free radical polymerization)
     9003-49-0P, Poly(butyl acrylate)
                                      9003-53-6P, Polystyrene
TΤ
        (new seven- and eight-membered cyclic alkoxyamines
        catalysts for living free radical polymerization)
ΙT
     585-71-7, (1-Bromoethyl) benzene
        (preparation of initiator; new seven- and eight-membered
        cyclic alkoxyamines catalysts for living free radical
       polymerization)
ΙT
     109-63-7, Boron trifluoride etherate
        (ring enlargement; new seven- and eight-membered cyclic
        alkoxyamines catalysts for living free radical polymerization)
ΙT
     18107-18-1, (Trimethylsilyl)diazomethane
        (ring enlargement; new seven- and eight-membered cyclic
        alkoxyamines catalysts for living free radical polymerization)
REFERENCE COUNT:
                         40
                               THERE ARE 40 CITED REFERENCES AVAILABLE FOR
                               THIS RECORD. ALL CITATIONS AVAILABLE IN THE
                               RE FORMAT
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L68 ANSWER 16 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2003:148870 HCAPLUS Full-text

DOCUMENT NUMBER: 139:22531

A convenient synthesis of functionalized TITLE:

alkoxyamines as initiators for living free radical

polymerization

Sugimoto, Naoya; Narumi, Atsushi; Satoh, AUTHOR(S):

Toshifumi; Kaga, Harumi; Kakuchi, Toyoji

Graduate School of Engineering, Division of CORPORATE SOURCE:

Molecular Chemistry, Hokkaido University, Sapporo,

060-8628, Japan

SOURCE: Polymer Bulletin (Berlin, Germany) (2003), 49(5),

337-340

CODEN: POBUDR; ISSN: 0170-0839

PUBLISHER: Springer-Verlag

DOCUMENT TYPE: Journal LANGUAGE: English Entered STN: 27 Feb 2003 ED

2,2,6,6-Tetramethylpiperidine-1-oxyl (TEMPO) was reacted with ethylbenzene (1a), 1-bromo-4-ethylbenzene (1b), and 4-ethylphenyl acetate (1c), resp., using tert-BuOOWCo(OAc)·4H2O in acetonitrile at room temperature The reactions produced the resp. TEMPO-adducts (2a, 2b, and 2c) in the yields of 37, 44, and 45%, which were based on TEMPO. Similarly, TEMPO was reacted with 4-ethylphenyl 2,3,6,2',3',4',6'-hepta-O-acetyl- $\beta$ -D-cellobioside (1d) to afford the glycoconjugated TEMPO-adduct (2d) in 45% yield, which was based on 1d. These results indicated that the reaction has the potential to become an easy and also safe strategy, which provided various functionalized alkoxylamines.

ΙT 154554-67-3P

> (synthesis of functionalized alkoxyamines as initiators for living free radical polymerization)

RN 154554-67-3 HCAPLUS

Piperidine, 2,2,6,6-tetramethyl-1-(1-phenylethoxy)- (CA INDEX NAME) CN

CC 35-3 (Chemistry of Synthetic High Polymers)

ST alkoxyamine initiator living radical polymn

154554-67-3P 178625-97-3P 213699-59-3P ΙT

538373-53-4P

(synthesis of functionalized alkoxyamines as initiators for living

free radical polymerization)

THERE ARE 10 CITED REFERENCES AVAILABLE FOR REFERENCE COUNT: 10

THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L68 ANSWER 17 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2003:140776 HCAPLUS Full-text

DOCUMENT NUMBER: 139:36837

TITLE: Nitroxide-mediated radical polymerization of

styrene in miniemulsion: model studies of

alkoxyamine-initiated systems

AUTHOR(S): Ma, John W.; Smith, Jodi A.; McAuley, Kim B.;

Cunningham, Michael F.; Keoshkerian, Barkev;

Georges, Michael K.

CORPORATE SOURCE: Department of Chemical Engineering, Queen's

University, Kingston, ON, K7L 3N6, Can.

SOURCE: Chemical Engineering Science (2003), 58(7),

1163-1176

CODEN: CESCAC; ISSN: 0009-2509

PUBLISHER: Elsevier Science Ltd.

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 25 Feb 2003

A math. model was developed to describe the behavior of the nitroxide-mediated AB miniemulsion polymerization (NMMP) of styrene initiated by alkoxyamine initiators. The model includes mechanisms describing reactions in the aqueous and organic phases, particle nucleation, the entry and exit of oligomeric radicals, and the partitioning of nitroxide and styrene between the aqueous and organic phases. The influence of nitroxide partitioning on the polymerization kinetics was examined by modeling systems initiated by the alkoxyamines BST and hydroxyl-BST; BST and hydroxyl-BST are benzoylstyryl radicals terminated by the nitroxides TEMPO and 4-hydroxyl-TEMPO, resp. Predicted monomer conversions, number average mol. wts. and polydispersities were in agreement with exptl. measured values. Simulations and math. anal. showed that the rate of styrene NMMP is not strongly influenced by the partitioning properties of TEMPO and 4-hydroxyl-TEMPO because of the complex interaction between reaction equilibrium, phase equilibrium, termination and thermal initiation. However, in the absence of styrene thermal initiation, nitroxide partitioning had a significant influence on the polymerization kinetics. The model was also used to make quant. ests. of: the population of active and dormant polymer radicals derived from both alkoxyamine initiators and thermal initiation; the population of dead polymer chains; and the number mol. weight distributions of living and dead polymer chains.

IT 81913-53-3

(polymerization catalyst; math. modeling of nitroxide-mediated radical polymerization of styrene in miniemulsion initiated by alkoxyamine initiators)

RN 81913-53-3 HCAPLUS

CN Benzeneethanol,  $\beta$ -[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]-, 1-benzoate (CA INDEX NAME)

CC 35-3 (Chemistry of Synthetic High Polymers)

ST math modeling living radical polymn styrene; simulation nitroxide mediated polymn styrene miniemulsion; alkoxyamine initiator living radical polymn styrene miniemulsion

IT Polymerization kinetics

(living, radical; math. modeling of nitroxide-mediated living

radical polymerization of styrene in miniemulsion initiated by alkozyamine initiators) ΙT Polymerization (living, radical; math. modeling of nitroxide-mediated radical polymerization of styrene in miniemulsion initiated by alkoxyamine ΙT Simulation and Modeling (math. modeling of nitroxide-mediated living radical polymerization of styrene in miniemulsion initiated by alkoxyamine Molecular weight ΤТ Polydispersity (math. modeling of nitroxide-mediated radical polymerization of styrene in miniemulsion initiated by alkoxyamine initiators ΙT 100-42-5, Styrene, reactions (math. modeling of nitroxide-mediated radical polymerization of styrene in miniemulsion initiated by alkoxyamine initiators 9003-53-6P, Polystyrene ΙT (math. modeling of nitroxide-mediated radical polymerization of styrene in miniemulsion initiated by alkoxyamine initiators 81913-53-3 470689-10-2 ΙT (polymerization catalyst; math. modeling of nitroxide-mediated radical polymerization of styrene in miniemulsion initiated by alkoxyamine initiators) REFERENCE COUNT: 47 THERE ARE 47 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L68 ANSWER 18 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2002:928481 HCAPLUS Full-text 138:137679 DOCUMENT NUMBER: n-Butyl acrylate polymerization mediated by a TITLE: PROXYL nitroxide AUTHOR(S): Cameron, Neil R.; Reid, Alistair J. Department of Chemistry, University of Durham, CORPORATE SOURCE: Durham, DH1 3LE, UK Macromolecules (2002), 35(27), 9890-9895 SOURCE: CODEN: MAMOBX; ISSN: 0024-9297 American Chemical Society PUBLISHER: DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 08 Dec 2002 N-Bu acrylate has been polymerized in the presence of an alkozyamine initiator AB derived from the PROXYL nitroxide 2,2',5-trimethyl-5'-phenylpyrrolidinyl-1oxyl. It was found that polymns. were rapid, reaching almost complete conversion within 2 h; in contrast, no conversion was observed when TEMPO was used as the mediator. The addition of a small amount of free nitroxide resulted in slower polymns. although an induction period, the length of which varied with excess nitroxide concentration, was observed Size exclusion chromatog. indicated that polymerization control was poor; Mn initially increased rapidly and then much more slowly, and polydispersities were found to be broad and to increase with conversion. Quant. 13C NMR spectroscopy revealed the resulting poly(Bu acrylate) to be branched. Despite the poor control, the PBA was able to act as a macroinitiator for the polymerization of styrene, yielding a block copolymer with a growth in Mn and a reduction of

polydispersity with conversion. It is suggested that the polymerization of Bu

acrylate in the presence of the substituted PROXYL derivative is living but not controlled.

IT 185055-65-6

(Bu acrylate polymerization mediated by PROXYL nitroxide)

RN 185055-65-6 HCAPLUS

CN Piperidine, 1-[2-(1,1-dimethylethoxy)-1-phenylethoxy]-2,2,6,6-tetramethyl- (CA INDEX NAME)

CC 35-4 (Chemistry of Synthetic High Polymers)

IT 62539-49-5 185055-65-6 328311-61-1

(Bu acrylate polymerization mediated by PROXYL nitroxide)

REFERENCE COUNT: 38 THERE ARE 38 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L68 ANSWER 19 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2001:886070 HCAPLUS Full-text

DOCUMENT NUMBER: 136:20022

TITLE: Process for the synthesis of amine ethers from

N-oxylamines and hydrocarbons in the presence of

hydroperoxides and copper catalysts.

INVENTOR(S): Hafner, Andreas; Kirner, Hans Juerg;

Schwarzenbach, Franz; Van Der Schaaf, Paul

Adriaan; Nesvadba, Peter

PATENT ASSIGNEE(S): Ciba Specialty Chemicals Holding Inc., Switz.

SOURCE: PCT Int. Appl., 84 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PAT	PATENT NO.				KIND DATE		APPLICATION NO.						DATE				
	2001						2001		1	WO 2	001-	EP56	68		20010517		
WO	2001	0922	28		A3		2002	0516									
	W:	ΑE,	ΑG,	ΑL,	ΑM,	ΑT,	ΑU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	BZ,	CA,	CH,	
		CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,	
		GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KP,	KR,	KΖ,	
		LC,	LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	
		NO, NZ, PL,		PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	ТJ,	TM,	TR,	
		TT,	TZ,	UA,	UG,	US,	UZ,	VN,	YU,	ZA,	ZW						
	RW:	GH,	GM,	KE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZW,	ΑT,	BE,	CH,	
		CY,	DE,	DK,	ES,	FI,	FR,	GB,	GR,	ΙE,	ΙΤ,	LU,	MC,	NL,	PT,	SE,	
		TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GW,	$\mathrm{ML}_{\prime}$	MR,	ΝE,	SN,	TD, TG	
TW	5728	96			В		2004	0121	1	TW 2	001-	9011	0396		2	0010501	
CA	A 2407866 A1						2001	1206	CA 2001-2407866						20010517		
EP	P 1284966 A2 20030226 EP 200					001-	9451.	52		2	0010517						
	R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	ΙΤ,	LI,	LU,	NL,	SE,	MC,	
	PT, IE, SI, LT, 1					LV,	FI,	RO,	O, MK, CY, AL, TR								

JP 2003535080	T	20031125	JP	2002-500843		20010517
RU 2273634	C2	20060410	RU	2002-133210		20010517
US 2003171461	A1	20030911	US	2002-296107		20021122
US 6900328	В2	20050531				
US 2005043552	A1	20050224	US	2004-919776		20040817
PRIORITY APPLN. INFO.:			EP	2000-810461	A	20000526
			WO	2001-EP5668	W	20010517
			US	2002-296107	A1	20021122

OTHER SOURCE(S): CASREACT 136:20022; MARPAT 136:20022

ED Entered STN: 07 Dec 2001

AB [G1G2T2CN(T1)0]aE1 [a = 1, 2; when a = 1, E = alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, (unsatd.) aliphatic bicyclic or tricyclic hydrocarbyl, aralkyl, etc.; when a = 2, E = alkylene, cycloalkylene, cycloalkenylene, alkenylene, phenylalkyl, etc.; T1 = (substituted) tertiary alkyl, Ph, cycloalkyl, heterocyclyl, etc.; T2 = H, halo, NO2, cyano, organic radical; T1T2 = organic linking group; G1, G2 = H, halo, NO2, cyano, aminocarbonyl, etc.], were prepared by reaction of G1G2T2CN(T1)O· with EH in the presence of an organic hydroperoxide and catalytic Cu or Cu compds. Thus, 1-oxyl-2,2,6,6-tetramethyl-4- propoxypiperidine, ethylbenzene, tert-Bu hydroperoxide in decane, and CuCl2 in EtOH were stirred at 60° for 60 min. to give 97% 1-(1-phenethyloxy)-2,2,6,6-tetramethyl-4-propoxypiperidine.

IT 154554-67-3P

(process for the synthesis of amine ethers from N-oxylamines and hydrocarbons in the presence of hydroperoxides and copper catalysts)

RN 154554-67-3 HCAPLUS

CN Piperidine, 2,2,6,6-tetramethyl-1-(1-phenylethoxy)- (CA INDEX NAME)

IC ICM C07D211-94

ICS C07D405-12; C07D401-12; C08K005-3435; C08F004-00; C07D217-26; C07D209-44; C07D241-08

CC 27-16 (Heterocyclic Compounds (One Hetero Atom)) Section cross-reference(s): 28, 35

ST amine ether prepn; oxylamine hydrocarbon coupling reaction hydroperoxide copper catalyst; piperidinoxyl hydrocarbon coupling reaction hydroperoxide copper catalyst; hydrocarbyloxyamine prepn heat light stabilizer; antioxidant hydrocarbyloxyamine prepn; polymn regulator hydrocarbyloxyamine prepn IT Coupling reaction

(homolytic; process for the synthesis of amine ethers from N-oxylamines and hydrocarbons in the presence of hydroperoxides and copper catalysts)

IT Phase transfer catalysts

(process for the synthesis of amine ethers from N-oxylamines and hydrocarbons in the presence of hydroperoxides and copper

catalysts)

IT 1643-19-2, Tetrabutylammonium bromide 7440-50-8, Copper, uses 7440-50-8D, Copper, compds. 7447-39-4, Cupric chloride, uses 7758-89-6, Cuprous chloride 7787-70-4, Cuprous bromide 35675-80-0, Trioctyl-methyl ammonium bromide

(process for the synthesis of amine ethers from N-oxylamines and hydrocarbons in the presence of hydroperoxides and copper catalysts)

IT 7803-49-8DP, Hydroxylamine, hydrocarbyloxyamines 154554-67-3P

243972-01-2P 335201-37-1P 378245-15-9P

378245-16-0P 378245-17-1P 378245-18-2P 378245-19-3P

378245-20-6P 378245-21-7P 378245-22-8P 378245-23-9P

378245-24-0P 378245-25-1P 378245-26-2P 378245-27-3P 378245-28-4P 378245-29-5P 378245-30-8P 378245-31-9P 378245-32-0P 378245-33-1P 378245-34-2P

(process for the synthesis of amine ethers from N-oxylamines and hydrocarbons in the presence of hydroperoxides and copper catalysts)

IT 100-41-4, Ethylbenzene, reactions 101-41-7, Methyl phenylacetate 103-45-7, 2-Phenethyl acetate 110-82-7, Cyclohexane, reactions 110-83-8, Cyclohexene, reactions 110-87-2, 3,4-Dihydro-2H-pyran 140-29-4, Benzyl cyanide 536-75-4, 4-Ethylpyridine 1746-13-0, Allyl phenyl ether 2226-96-2, 4-Hydroxy-2,2,6,6-tetramethylpiperidin-1-oxyl 2564-83-2, 2,2,6,6-Tetramethylpiperidin-1-oxyl 2930-02-1 3282-30-2, Pivaloyl chloride 14691-89-5, 4-(Acetylamino)-2,2,6,6-tetramethylpiperidin-1-oxyl 18554-09-1 61746-17-6 71809-12-6 104134-81-8 154186-10-4 194147-18-7 264279-93-8 (process for the synthesis of amine ethers from N-oxylamines and hydrocarbons in the presence of hydroperoxides and copper

TT 75-98-9P, 2,2-Dimethylpropionic acid 182190-87-0P 378245-35-3P 378245-36-4P 378245-37-5P (process for the synthesis of amine ethers from N-oxylamines and hydrocarbons in the presence of hydroperoxides and copper

catalysts)

IT 75-91-2, tert-Butyl hydroperoxide 79-21-0, Peracetic acid
(process for the synthesis of amine ethers from N-oxylamines and
hydrocarbons in the presence of hydroperoxides and copper
catalysts)

L68 ANSWER 20 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2001:868459 HCAPLUS <u>Full-text</u>

DOCUMENT NUMBER: 136:6539

catalysts)

TITLE: Hydroxylamine esters as polymerization

initiators

INVENTOR(S): Roth, Michael; Pfaendner, Rudolf; Nesvadba, Peter;

Zink, Marie-Odile

PATENT ASSIGNEE(S): Ciba Specialty Chemicals Holding Inc., Switz.

SOURCE: PCT Int. Appl., 114 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

----WO 2001090113 A1 20011129 WO 2001-EP5447 20010514
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH,

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CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD,
             GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ,
             LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,
            NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR,
             TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW
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                                          BR 2001-10854
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                                20030211
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     EP 1282630
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                         В1
                                20060510
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                                20031118 JP 2001-586300
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     JP 2003534347
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                               20060510
                                          EP 2006-100223
                                                                   20010514
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     PT 1282630
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    ES 2262647
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     RU 2298563
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                                          RU 2002-133442
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     TW 249539
                              20060221
                                          TW 2001-90111569
                                                                  20010515
                        B1 20071207 KR 2002-714001
     KR 783605
                                                                  20021018
    US 2003216494
                       A1 20031120 US 2002-275495
B2 20060418
                                                                   20021105
     US 7030196
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                                                                  20021113
                       A 20030106 NO 2002-5532
A 20030425 MX 2002-PA11344
A 20031016 ZA 2002-9397
     NO 2002005532
                                                                   20021118
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                                                                   20021118
     ZA 2002009397
                                                                   20021119
    US 2006128903 A1 20060615 US 2006-339214 IN 2007CN01321 A 20070831 IN 2007-CN1321 KR 2007086650 A 20070827 KR 2007-714491
                                                                   20060125
                                            IN 2007-CN1321
                                                                   20070330
                                           KR 2007-714491 20070625
EP 2000-810443 A 20000519
PRIORITY APPLN. INFO.:
                                            EP 2001-931694 A3 20010514
                                            WO 2001-EP5447 W 20010514
                                            KR 2002-714001 A3 20021018
                                            US 2002-275495
                                                               A1 20021105
                                            IN 2002-CN1865
                                                              A3 20021113
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OTHER SOURCE(S): MARPAT 136:6539

ED Entered STN: 30 Nov 2001

AB The invention relates to novel cyclic and open-chain hydroxylamine esters and polymerizable compns. comprising these hydroxylamine esters and an ethylenically unsatd. monomer or oligomer. The invention also relates to use as polymerization initiators and to the use of known hydroxylamine esters and the novel hydroxylamine esters for the controlled degradation of polypropylene and for achieving a controlled increase in the mol. weight of polyethylene.

IT 7031-92-7P

(hydroxylamine esters as polymerization initiators and controlling degradation and mol. weight of polymers)

RN 7031-92-7 HCAPLUS

CN 4-Piperidinol, 1-(benzoyloxy)-2,2,6,6-tetramethyl-, benzoate (ester) (8CI, 9CI) (CA INDEX NAME)

ICM C07F009-59

IC

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ICS C07D295-24; C07D211-94; C07D241-08; C08F004-00; C08F008-50
CC
    35-3 (Chemistry of Synthetic High Polymers)
    Section cross-reference(s): 27, 67
ST
    hydroxylamine ester manuf polymn catalyst; radical polymn
    initiator hydroxylamine ester manuf
    Molecular weight
ΙT
    Polymer degradation
    Polymer degradation catalysts
        (hydroxylamine esters as polymerization initiators and
        controlling degradation and mol. weight of polymers)
    EPDM rubber
ΙT
    Linear low density polyethylenes
        (hydroxylamine esters as polymerization initiators and
        controlling degradation and mol. weight of polymers)
ΙT
    Polymer blends
        (hydroxylamine esters as polymerization initiators and
        controlling degradation and mol. weight of polymers)
ΙT
    Polymerization catalysts
        (radical; hydroxylamine esters as polymerization initiators and
        controlling degradation and mol. weight of polymers)
ΙT
    9002-88-4, Polyethylene
        (Lupolen 1812E, Hostalen GB 7250; hydroxylamine esters as polymerization
        initiators and controlling degradation and mol. weight of
       polymers)
ΙT
    376587-77-8P
       (ester of acetic acid; hydroxylamine esters as polymerization
       initiators and controlling degradation and mol. weight of
       polymers)
ΙT
    7031-92-7P 89108-37-2P 99365-17-0P
                                           113682-53-4P
    122809-58-9P 123978-94-9P 130048-66-7P 271242-59-2P
    376587-78-9P
                   376587-79-0P
                                  376587-80-3P
                                                 376587-81-4P
    376587-82-5P
                   376587-83-6P 376587-84-7P
                                                 376587-85-8P
    376587-86-9P 376587-87-0P 376587-88-1P 376587-89-2P
    376587 - 90 - 5P 376587 - 91 - 6P 376587 - 92 - 7P 376587 - 93 - 8P
    376587-94-9P 376587-95-0P 376587-96-1P 376587-97-2P
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    376588-03-3P
                   376588-04-4P
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    376588-11-3P 376588-13-5P 376588-15-7P
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    376588-19-1P 376588-20-4P 376588-21-5P 376588-24-8P
    376588-27-1P 376588-29-3P 376588-30-6P 376588-31-7P
    376588-32-8P 376588-33-9P 376588-34-0P 376588-36-2P
    376588-37-3P 376588-38-4P 376588-39-5P 376588-40-8P
    376588-41-9P
                   376588-42-0P
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(hydroxylamine esters as polymerization initiators and controlling degradation and mol. weight of polymers) ΙT 376588-00-0P (hydroxylamine esters as polymerization initiators and controlling degradation and mol. weight of polymers) 9003-49-0P, Butyl acrylate homopolymer ΙT (hydroxylamine esters as polymerization initiators and controlling degradation and mol. weight of polymers) 376588-35-1P ΙT 199101-44-5P (hydroxylamine esters as polymerization initiators and controlling degradation and mol. weight of polymers) ΙT 74-85-1D, Ethylene, polymers with  $\alpha$ -olefins 9003-07-0, Polypropylene 25085-53-4, Profax 6501 (hydroxylamine esters as polymerization initiators and controlling degradation and mol. weight of polymers) 79-21-0, Peracetic acid 79-37-8, Oxalyl chloride 98-88-4, Benzoyl ΙT chloride 102-92-1, Cinnamoyl chloride 103-71-9, Phenyl isocyanate, reactions 108-24-7, Acetic anhydride 109-90-0, Ethyl isocyanate 111-50-2, Adipoyl chloride 112-76-5, Stearoyl chloride 134-03-2, Sodium ascorbate 407-25-0, Trifluoroacetic anhydride 938-18-1, 2,4,6-Trimethylbenzovl chloride 1310-73-2, Sodium hydroxide, reactions 1499-21-4, Diphenylphosphinic chloride 2094-72-6, 1-Adamantylcarboxylic acid chloride 2516-91-8 2516-92-9 3282-30-2, Pivaloyl chloride 3637-10-3, 1,4-Dihydroxy-2,2,6,6tetramethylpiperidine 4972-13-8 6599-87-7 7803-49-8, Hydroxylamine, reactions 14691-89-5 16256-42-1, 2, 2, 5, 5-Tetramethyl-4-oxoimidazolidine 22977-67-9 24424-99-5 51210-48-1 61682-93-7 61745-37-7 61746-17-6 61964-43-0 71981-32-3 80037-90-7 98254-32-1 150981-00-3 151419-17-9 154186-10-4 195300-91-5, Chimassorb 2020 244020-99-3 244021-06-5 258289-25-7 264279-93-8 264280-22-0 376588-12-4 376588-14-6 376588-16-8 (hydroxylamine esters as polymerization initiators and controlling degradation and mol. weight of polymers) REFERENCE COUNT: 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L68 ANSWER 21 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2001:718062 HCAPLUS Full-text DOCUMENT NUMBER: 135:273360 Vinyl group-containing alkoxyamines, manufacture TITLE: of them and radical polymerization initiators, and graft polymers manufactured using the initiators INVENTOR(S): Hayashi, Masaki; Nakamura, Tomoyuki; Ujigawa, Norihisa NOf Corporation, Japan PATENT ASSIGNEE(S): SOURCE: Jpn. Kokai Tokkyo Koho, 20 pp. CODEN: JKXXAF DOCUMENT TYPE: Patent LANGUAGE: Japanese FAMILY ACC. NUM. COUNT: 1 PATENT INFORMATION: PATENT NO. KIND DATE APPLICATION NO. \_\_\_\_ \_\_\_\_\_\_ JP 2000-125936 20000426 JP 2000-8340 A 20000117 A 20011002 JP 2001270859 PRIORITY APPLN. INFO.:

OTHER SOURCE(S): MARPAT 135:273360

ED Entered STN: 02 Oct 2001

AB R1CO2CH2CR8R9ON(CR2R3R4)CR5R6R7 [R1 = monovalent organic group having radically polymerizable vinyl group; R2-R7 = C1-4 linear or branched alkyl; R2R7 may form (un)substituted C4-12 ring; R8 = H, Me; R9 = Ph, MeC6H4, cyano, C1-4 alkoxycarbonyl, C1-4 alkylcarbonyloxy] are manufactured by decomposition of (R1CO2)2 (R1 = same as above) in the presence of ·ON(CR2R3R4)CR5R6R7 (R2-R7 = same as above) and H2C:CR8R9 (R8, R9 = same as above). Thus, styrene was reacted with 2,2,6,6-tetramethylpiperidine-1-oxyl and cinnamoyl peroxide to give PhCH:CHCO2CH2CHPhOZ (Z = 2,2,6,6-tetramethylpiperidino), which was polymerized with (chloromethyl)styrene to give alkoxyamine-modified poly[(chloromethyl)styrene] with Mn 48,000. Styrene was polymerized in the presence of the initiator to give (chloromethyl)styrene-styrene graft copolymer.

IT 362660-50-2P

 $\hbox{(manufacture of vinyl group-containing alkoxyamines for manufacture of polymerization}\\$ 

initiators and graft polymers)

RN 362660-50-2 HCAPLUS

CN 2-Propenoic acid, 3-phenyl-, 2-phenyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl ester, (2E)- (CA INDEX NAME)

Double bond geometry as shown.

IC ICM C07C239-20

ICS C07D211-94; C08F004-00; C08F216-14; C08F218-00; C08F220-36; C08F291-00

CC 35-3 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 27

IT 362660-50-2P 362660-51-3P 362660-52-4P 362660-53-5P 362660-54-6P 362660-55-7P 362660-56-8P 362660-57-9P 362660-58-0P

(manufacture of vinyl group-containing alkoxyamines for manufacture of polymerization

initiators and graft polymers)

IT 362660-59-1P 362660-60-4P 362660-61-5P

(polymerization initiator; manufacture of vinyl group-containing alkoxyamines for

manufacture of polymerization initiators and graft polymers)

L68 ANSWER 22 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2001:662112 HCAPLUS Full-text

DOCUMENT NUMBER: 135:372063

TITLE: Synthesis of poly(methylene-b-styrene) by

sequential living polymerization Zhou, Xian-Zhi; Shea, Kenneth J.

AUTHOR(S): Zhou, Xian-Zhi; Shea, Kenneth J. CORPORATE SOURCE: Department of Chemistry, University of California,

Irvine, CA, 92696-2025, USA

SOURCE: Polymer Preprints (American Chemical Society,

Division of Polymer Chemistry) (2001), 42(2),

411-412

CODEN: ACPPAY; ISSN: 0032-3934

PUBLISHER: American Chemical Society, Division of Polymer

Chemistry

DOCUMENT TYPE: Journal; (computer optical disk)

LANGUAGE: English ED Entered STN: 11 Sep 2001

AB We have recently developed a synthesis of telechelic polymethylene. Organoboranes serve as the initiator and dimethylsulfoxonium methylide (1) functions as the monomer. The carbon chain is built up one carbon at a time by repetitive homologation (polyhomologation) of alkyl substituents. The living nature of the polyhomologation reaction permits control of the mol. weight as well as the functionality at either end of the polymer chain. We are utilizing polyhomologation chemical to develop general methods for the synthesis of well defined copolymers of polymethylene and the major commodity polyolefins. AB block copolymer of poly(methylene-b-styrene) was chosen as the targets of this research. The block copolymers are useful compatibilizers in PE/PS polymer blends.

IT 81913-53-3P

(polymerization initiator synthesis; synthesis of poly(methylene-b-styrene) by sequential living polymerization)

RN 81913-53-3 HCAPLUS

CN Benzeneethanol,  $\beta$ -[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]-, 1-benzoate (CA INDEX NAME)

CC 35-4 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 37

IT 94-36-0, Benzoyl peroxide, reactions 100-42-5, Styrene, reactions 2564-83-2, TEMPO

(polymerization initiator synthesis; synthesis of
poly(methylene-b-styrene) by sequential living polymerization)

IT 81913-53-3P

(polymerization initiator synthesis; synthesis of

poly(methylene-b-styrene) by sequential living polymerization)

IT 161776-41-6P

(polymerization initiator; synthesis of poly(methylene-b-

styrene) by sequential living polymerization)

REFERENCE COUNT: 23 THERE ARE 23 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L68 ANSWER 23 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2001:582410 HCAPLUS <u>Full-text</u>

DOCUMENT NUMBER: 135:289127

TITLE: Nitroxide-Mediated Miniemulsion Acrylate

Polvmerization

AUTHOR(S): Keoshkerian, Barkev; Szkurhan, Andrea R.; Georges,

Michael K.

CORPORATE SOURCE: Xerox Research Centre of Canada, Mississauga, ON,

L5K 2L1, Can.

SOURCE: Macromolecules (2001), 34(19), 6531-6532

CODEN: MAMOBX; ISSN: 0024-9297

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 12 Aug 2001

AB Bu acrylate can be polymerized by a stable free radical polymerization miniemulsion process at 135° using alkoxyamine terminated with TEMPO as initiator in the presence of a small amount of ascorbic acid. The conversion was 60-65%. Polydispersities were broader than when hydroxy-containing nitroxide was used, but the polymers were living as seen by the incremental

increase in mol. weight with time.

IT 81913-53-3

(miniemulsion polymerization of Bu acrylate in presence of)

RN 81913-53-3 HCAPLUS

CN Benzeneethanol,  $\beta$ -[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]-, 1-benzoate (CA INDEX NAME)

CC 35-4 (Chemistry of Synthetic High Polymers)

IT Polymerization catalysts

(emulsion; miniemulsion polymerization of Bu acrylate in presence of TEMPO-terminated alkoxyamine initiator)

IT 81913-53-3

(miniemulsion polymerization of Bu acrylate in presence of)

IT 9003-49-0P, Poly(butyl acrylate)

(miniemulsion polymerization of Bu acrylate in presence of

TEMPO-terminated alkoxyamine initiator)

REFERENCE COUNT: 16 THERE ARE 16 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L68 ANSWER 24 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2001:165813 HCAPLUS Full-text

DOCUMENT NUMBER: 134:223147

TITLE: Functionalized alkoxyamine

initiators for radical polymerization and

their preparation

INVENTOR(S): Melchiors, Martin; Hoecker, Hartwig; Keul, Helmut;

Achten, Dirk

PATENT ASSIGNEE(S): Bayer A.-G., Germany SOURCE: Ger. Offen., 13 pp.

CODEN: GWXXBX

DOCUMENT TYPE: Patent LANGUAGE: German

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

-	PATENT NO.			KIND DATE		APPLICATION NO.						DATE						
		1994				A1			20010308									990907
	EΡ	1083	169			A1		2001	0314	EI	Ρ	2000-	1181	24			20	000828
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			PT,	ΙE,	SI,	LT,	LV,	FI,	RO									
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	ES	2208				,				ES	S	2000-	1181	24			20	000828
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		6632						2003				2000-						000831
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		2317				A1		2001				2000-						000901
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		2001				A		2001				2000-						000907
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		6800				В2		2004	1005		_					_		
PRIOR	ILX	APP	LN.	INFO	.:					DE	Ε	1999-	1994	2614		А	19	990907
										DI	Ε	1999-	1994	2615		Α	19	990907
										US	S	2000-	6520	75		А3	20	000831

OTHER SOURCE(S): MARPAT 134:223147

ED Entered STN: 09 Mar 2001

GΙ

AB Alkoxyamines HOCHR1CR2R3ONR4R5 (I) (more specifically, II are claimed as new compds.) [R1-R3 = H, C1-20 (cyclo)alkyl, C6-24 aryl, halogen, CN, amide or ester group; R4, R5 = C1-24 organic group in which the N-linked C bears no H; R4 + R5 optionally contains ≥1 reactive functional group; R6 = H, C1-20 (cyclo)alkyl; R8-R11 = C1-20 (cyclo)alkyl, C6-24 aryl, (un)substituted C7-C24 aralkyl, ring portion; R17 = H, reactive functional group; R18 = C1-20 (cyclo)alkyl] are prepared by reaction of R1CH:CR2R3 with R4R5NO in the presence of H2O2 and a reducing agent, and are used as initiators of radical polymerization of vinyl monomers. Thus, addition of 50 mL 30% H2O2 dropwise over 5 h to a solution of 1 mol styrene and 0.1 mol 2,2,6,6-tetramethyl-1-piperidinyloxy in 300 mL MeOH at 40° containing 0.12 mol FeSO4.7H2O and 0.24 mol NaHCO3 gave I [R1 = R2 = H, R3 = Ph, R4R5 = CMe2(CH2)3CMe2] in 71% yield. IT 161776-41-6P

(preparation of functionalized alkoxyamine initiators for radical polymerization)

RN 161776-41-6 HCAPLUS

Benzeneethanol,  $\beta$ -[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]- (CA CN INDEX NAME)

IC ICM C07D211-94

ICS C07D247-00; C07D273-00

35-3 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 27

ST alkoxyamine initiator vinyl polymn

Polymerization catalysts ΙT

(preparation of functionalized alkoxyamine initiators

for radical polymerization)

161776-41-6P 183959-05-9P 329180-72-5P ΤT 329326-50-3P 329326-51-4P 329326-52-5P

329326-53-6P

(preparation of functionalized alkoxyamine initiators for radical polymerization)

329326-49-0P ΤТ

> (preparation of functionalized alkoxyamine initiators for radical polymerization)

ΙT 144-55-8, Sodium bicarbonate, uses

(preparation of functionalized alkoxyamine initiators

for radical polymerization)

96-33-3, Methyl acrylate ΙT 80-62-6, Methyl methacrylate Butyl methacrylate 100-42-5, Styrene, reactions 101-43-9, Cyclohexyl methacrylate 103-11-7, 2-Ethylhexyl acrylate 108-31-6, Maleic anhydride, reactions 141-32-2, Butyl acrylate 2226-96-2, 1-Piperidinyloxy, 4-hydroxy-2,2,6,6-tetramethyl- 2564-83-2, TEMPO 7534-94-3, Isobornyl methacrylate 7722-84-1, Hydrogen peroxide, reactions 14691-89-5

> (preparation of functionalized alkoxyamine initiators for radical polymerization)

7720-78-7, Ferrous sulfate ΤТ

> (reducing agent; preparation of functionalized alkoxyamine initiators for radical polymerization)

L68 ANSWER 25 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2001:165812 HCAPLUS Full-text

DOCUMENT NUMBER: 134:223167

TITLE: Process for the manufacture of telechelic vinyl

oligomers and polymers and their use

INVENTOR(S): Melchiors, Martin; Hoecker, Hartwig; Keul, Helmut;

Achten, Dirk

PATENT ASSIGNEE(S): Bayer A.-G., Germany SOURCE: Ger. Offen., 12 pp.

CODEN: GWXXBX

DOCUMENT TYPE: Patent LANGUAGE: German

FAMILY ACC. NUM. COUNT: 2 PATENT INFORMATION:

PA.	ATENT NO.			KIND DATE		APPLICATION NO.						DATE				
DE	1994	 2614			A1	2	2001	0308	DE	19	999-	 1994	2614		-	19990907
EP	1083	169			A1	2	2001	0314	EP	20	0.00 - 1	1181	24			20000828
EP	1083	169			В1	2	2003	1029								
	R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB, G	R,	IT,	LI,	LU,	NL,	SE	E, MC,
		PT,	ΙE,	SI,	LT,	LV,	FΙ,	RO								
EP	1083	186			A1	2	2001	0314	EP	20	000-	1181	25			20000828
	R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB, G	R,	ΙT,	LI,	LU,	NL,	SI	E, MC,
		PT,	ΙE,	SI,	LT,	LV,	FΙ,	RO								
ES	2208	197			Т3	2	20040	0616	ES	20	000-	1181	24			20000828
US	6573	346			В1	2	2003	0603	US	20	000-	5520	75			20000831
US	6632	895			В1	2	2003	1014	US	20	000-	5522	03			20000831
CA	2317	321			A1	2	2001	0307	CA	20	000-	2317	321			20000901
CA	2317	323			A1	2	2001	0307	CA	20	000-	2317	323			20000901
JP	2001	0811	17		Α	2	2001	0327	JP	20	000-	2715	41			20000907
JP	2001	1067	13		Α	2	2001	0417	JP	20	000-	2715	49			20000907
US	2003	2080.	21		A1	2	2003	1106	US	20	003-	4086	25			20030407
US	6800	708			В2	2	20041	1005								
PRIORIT	Y APP	LN.	INFO	.:					DE	19	999-	1994	2614		A	19990907
									DE	19	99-	1994	2615		A	19990907
									US	20	000-	6520	75		А3	20000831

OTHER SOURCE(S): MARPAT 134:223167

ED Entered STN: 09 Mar 2001

The telechelic entities, of number-average mol. weight 500-10,000 and with OH at one chain end and a group reactive toward NCO, OH, CO2H, anhydride, or epoxy groups at the other, are prepared by polymerization of vinyl monomers in the presence of an alkozyamine initiator HOCHR1CR2R3ONR4R5 [R1-R3 = H, C1-20 (cyclo)alkyl, C6-24 aryl, halogen, CN, ester or amide group; R4, R5 = C1-24 organic group (at least partly aliphatic)]. They can then react with unsatd. alcs., isocyanates, acids, anhydrides or epoxides to form, e.g., macromonomers with OH functionality at the far end of the chain. Thus, 40 equiv styrene was polymerized for 4 h at 130° in the presence of 1 equiv 1-(2-hydroxy-1-phenylethoxy)-2,2,6,6-tetramethyl-4-piperidinol to give a polystyrene with Mn 2190 and OH functionality >1.9 in 53% yield.

IT 161776-41-6

(initiator in manufacture of telechelic vinyl oligomers and polymers)

RN 161776-41-6 HCAPLUS

CN Benzeneethanol,  $\beta$ -[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]- (CA INDEX NAME)

ICS C08F022-06; C08F012-08; C08F002-42

CC 35-4 (Chemistry of Synthetic High Polymers)

ST telechelic vinyl oligomer manuf; alkoxyamine

initiator vinyl monomer polymn

IT Polymers, preparation

(telechelic; manufacture of telechelic vinyl oligomers and polymers by use of alkoxyamine initiator)

IT 161776-41-6 183959-05-9 329180-72-5

(initiator in manufacture of telechelic vinyl oligomers and polymers)

L68 ANSWER 26 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2001:137295 HCAPLUS <u>Full-text</u>

DOCUMENT NUMBER: 134:179364

TITLE: Polymer blends with improved impact resistance INVENTOR(S): Chin, Hui; Botkin, James Harold; Fuso, Francesco;

Wunderlich, Wiebke

PATENT ASSIGNEE(S): Ciba Specialty Chemicals Holdings Inc., Switz.

SOURCE: PCT Int. Appl., 53 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

			KIND DATE			APPLICATION NO.						DATE					
							2001	0222	,	wo .	2000-	EP75	28			 20000	803
	W:	ΑE,	AG,	AL,	AM,	AT,	ΑU,	ΑZ,	BA,	BB	, BG,	BR,	BY,	BZ,	CA	, СН,	
		CN,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EE	, ES,	FΙ,	GB,	GD,	GE	, GH,	
		GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE	, KG,	ΚP,	KR,	KZ,	LC	, LK,	
		LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK	, MN,	MW,	MX,	MZ,	ИО	, NZ,	
		PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK	, SL,	ТJ,	TM,	TR,	ΤТ	, TZ,	
		UA,	UG,	UΖ,	VN,	YU,	ZA,	ZW,	ΑM,	ΑZ	, BY,	KG,	KΖ,	MD,	RU	, TJ,	TM
	RW:	GH,	GM,	ΚE,	LS,	MW,	MZ,	SD,	SL,	SZ	, TZ,	UG,	ZW,	ΑT,	BE	, СН,	
		CY,	DE,	DK,	ES,	FI,	FR,	GB,	GR,	IE	, IT,	LU,	MC,	NL,	PΤ	, SE,	
		BF,	ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	GW	, ML,	MR,	NE,	SN,	TD	, TG	
CA	2376.	569			A1		2001	0222	1	CA .	2000-	2376	569			20000	803
EP	1203	054			A1		2002	0508		EP .	2000-	9494	48			20000	803
EP	1203																
	R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR	, IT,	LI,	LU,	NL,	SE	, MC,	
		•	,	,	•	,	,	,	,		, AL						
											2001-						
											2000-						
ES	2231	225			Т3		2005	0516		ES .	2000-	9494	48			20000	803
	6444	_									2000-						
										US .	2002-	1920	91			20020	710
US	6765	062			В2		2004	0720									
PRIORIT	Y APP:	LN.	INFO	. :						US	1999-	1484	99P		P	19990	812
									,	WO .	2000-	EP75	28	,	W	20000	803
										US .	2000-	6344	41		A3	20000	808

ED Entered STN: 25 Feb 2001

AB Polymer blends, in particular polyphenylene ether-polyamide blends, are effectively compatibilized by incorporating certain oligomers, cooligomers, polymers and copolymers of narrow mol. weight distribution that are prepared under free radical polymerization conditions with glycidyl-functionalized nitroxyl initiators. The oligomers, cooligomers, polymers and copolymers contain ≥1 oxyamine group and ≥1 glycidyl-containing initiator group. Thus, a

blend of HPP 820 37.5, nylon 66 37.5, Kraton FG 10, and polystyrene having glycidyl and oxyamine groups 15 parts was kneaded/extruded, and injection molded into samples having notched Izod impact strength 1.67, tensile strength 5.0 kg/mm2, elongation 27%, and break energy 57 kg-mm; vs. 0.54, 6.2, 15, and 27, resp., for a 50:50 HPP 820:nylon 66 blend.

IT 243972-01-2P

(polymer blends with improved impact resistance)

RN 243972-01-2 HCAPLUS

CN Piperidine, 2,2,6,6-tetramethyl-1-[1-[4-(oxiranylmethoxy)phenyl]ethoxy ]-4-propoxy- (9CI) (CA INDEX NAME)

IC ICM C08L071-12

ICS C08L077-00; C08L067-00; C08L025-04

CC 37-6 (Plastics Manufacture and Processing)

IT 243972-01-2P

(polymer blends with improved impact resistance)

REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L68 ANSWER 27 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2001:61904 HCAPLUS Full-text

DOCUMENT NUMBER: 134:252044

TITLE: Factors Influencing the C-O Bond Homolysis of

Alkoxyamines: Effects of H-Bonding and Polar

Substituents

AUTHOR(S): Marque, Sylvain; Fischer, Hanns; Baier, Elisabeth;

Studer, Armido

CORPORATE SOURCE: Physikalisch-Chemisches Institut, Universitaet

Zuerich, Zurich, CH-8057, Switz.

SOURCE: Journal of Organic Chemistry (2001), 66(4),

1146-1156

CODEN: JOCEAH; ISSN: 0022-3263

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 134:252044

ED Entered STN: 26 Jan 2001

The synthesis of various new trialkylhydroxylamines is described. The rate constant of the C-O bond cleavage of these new alkoxyamines has been measured. For example, C-O bond homolysis rates in a series of para-substituted TEMPO-styryl compds. TEMPO-CH(CH3)C6H5X (p-MeO, p-Me, p-H, p-Br, and p-MeO2C) are presented. Furthermore, rate consts. for the C-O bond cleavage of  $\alpha$ -heteroaryl-substituted secondary alkoxyamines are discussed. A correlation by which the rate constant for the C-O bond cleavage of TEMPO-derived alkoxyamines can be predicted from the C-H BDEs of the corresponding alkanes is presented. Solvent effects as well as the effect of camphorsulfonic acid on the rate of the C-O bond homolysis are discussed. Finally, EPR and kinetic evidence show that alkoxyamines derived from nitroxides which are capable of

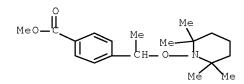
intramol. H-bonding undergo C-O bond cleavage faster than the corresponding non-H-bond-forming analogs.

IT 212132-39-3P

(effects of H-bonding and polar substituents on the C-O bond homolysis kinetics of alkoxyamines)

RN 212132-39-3 HCAPLUS

CN Benzoic acid, 4-[1-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl]-, methyl ester (CA INDEX NAME)



CC 22-8 (Physical Organic Chemistry)

Section cross-reference(s): 35

IT 212132-39-3P 270901-39-8P 270901-51-4P

270901-52-5P 270901-53-6P 270901-54-7P 270901-56-9P

270901-82-1P 270901-83-2P 270901-86-5P 270901-97-8P

270901-98-9P 270902-02-8P 330938-04-0P 330938-05-1P

330938-06-2P 330938-07-3P 330938-08-4P 330938-09-5P

330938-10-8P 330938-11-9P 330938-14-2P

(effects of H-bonding and polar substituents on the C-O bond

homolysis kinetics of alkoxyamines)

REFERENCE COUNT: 77 THERE ARE 77 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L68 ANSWER 28 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2000:911209 HCAPLUS <u>Full-text</u>

DOCUMENT NUMBER: 134:72025

TITLE: Preparation of vinyl-containing alkoxyamines used

as polymerizable monomers of polymeric radical initiators for manufacture of graft copolymers

INVENTOR(S): Hayashi, Masaki; Nakamura, Tomoyuki; Ujikawa,

Norihisa

PATENT ASSIGNEE(S): NOF Corporation, Japan SOURCE: PCT Int. Appl., 69 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.						KIN:	D	DATE		APPLICATION NO.						DATE		
-							_											
V	VΟ	2000	7870	09		A1		2000	1228	,	WO 2	000-	JP31	70		20000517		
		W:	ΑE,	AG,	AL,	AM,	ΑT,	AU,	AZ,	BA,	BB,	BG,	BR,	BY,	CA,	CH,	CN,	
			CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EE,	ES,	FI,	GB,	GD,	GE,	GH,	GM,	
			HR,	HU,	ID,	IL,	IN,	IS,	KE,	KG,	KP,	KR,	KΖ,	LC,	LK,	LR,	LS,	
			LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	NO,	NΖ,	PL,	PT,	RO,	
			RU,	SD,	SE,	SG,	SI,	SK,	SL,	ΤJ,	TM,	TR,	TT,	TZ,	UA,	UG,	US,	
			UZ,	VN,	YU,	ZA,	ZW											
		RW:	GH,	GM,	KE,	LS,	MW,	SD,	SL,	SZ,	TZ,	UG,	ZW,	ΑT,	BE,	CH,	CY,	
			DE,	DK,	ES,	FI,	FR,	GB,	GR,	IE,	IT,	LU,	MC,	NL,	PT,	SE,	BF,	

BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG JP 2000095744 Α 20000404 JP 1999-177422 19990623 JP 2001064308 Α 20010313 JP 1999-242052 19990827 AU 2000046142 Α 20010109 AU 2000-46142 20000517 EP 2000-927796 EP 1188742 20020320 20000517 Α1 EP 1188742 В1 20041215 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO Τ 20050115 AT 2000-927796 20000517 AT 284862 US 6919481 В1 20050719 US 2001-959754 20000517 PRIORITY APPLN. INFO.: JP 1999-177422 A 19990623 JP 1999-242052 A 19990827 JP 1998-206146 A 19980722 WO 2000-JP3170 W 20000517

OTHER SOURCE(S): MARPAT 134:72025

ED Entered STN: 29 Dec 2000

AB The title alkoxyamine (I) represented by general formula: R1COOCH2CH(C6H4CH:CH2)ONR3R4, wherein R1=aryl or R2O, R2=C1-12 linear or branched (cyclo)alkyl, R3, R4=C4-6 tertiary alkyl, or a C8-16 cyclic structure by bonding R3 and R4 together, is prepared by reacting a divinylbenzene with a nitroxide compound and an organic peroxide (II) at the decomposition temperature of II. Thus, heating the mixture of DVB-960 (mixture of m- and p-divinylbenzene) 50, 4-hydroxy-TEMPO 8.6 and diisopropyl peroxydicarbonate 11.6 g at 50° for 5 h gave an I, 0.62 g of which was mixed with 4.41 g Me methacrylate and 0.01 g tert-hexyl peroxypivalate and polymerized at 60° for 6 h to give a copolymer having graftable radical initiator sites.

IT 313975-54-1P

 $\hbox{(preparation of vinyl-containing alkoxyamines used as polymerizable } \\ \hbox{monomers}$ 

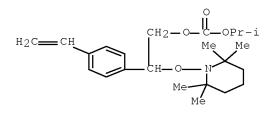
of polymeric radical initiators for manufacture of graft copolymers) 313975-54-1 HCAPLUS

RN 313975-54-1 HCAPLUS CN Carbonic acid, 2-(4-ethenylphenyl

Carbonic acid, 2-(4-ethenylphenyl)-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl 1-methylethyl ester, polymer with 1-(chloromethyl)-4-ethenylbenzene (9CI) (CA INDEX NAME)

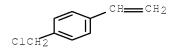
CM 1

CRN 262616-04-6 CMF C23 H35 N O4



CM 2

CRN 1592-20-7 CMF C9 H9 C1



IC ICM C07C239-20

ICS C07D211-94; C08F257-00; C08F004-00; C08F012-14

CC 35-2 (Chemistry of Synthetic High Polymers)

IT 313975-54-1P 313975-55-2P 314774-28-2P

314774-29-3P 314776-49-3P 314776-50-6P 314776-51-7P

314776-52-8P 314776-53-9P

 $\hbox{ (preparation of vinyl-containing alkoxyamines used as polymerizable } \\ \hbox{ monomers}$ 

of polymeric radical initiators for manufacture of graft copolymers)

IT 262616-03-5P 262616-04-6P 262616-05-7P

262616-06-8P 262616-07-9P 262616-08-0P

262616-09-1P 262616-10-4P 262616-11-5P

262616-12-6P 262846-44-6P 262846-45-7P 262846-46-8P

(radical-initiatable monomer; preparation of vinyl-containing alkoxyamines used as polymerizable monomers of polymeric radical initiators for

manufacture of graft copolymers)

REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L68 ANSWER 29 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2000:610052 HCAPLUS Full-text

DOCUMENT NUMBER: 133:322184

TITLE: Synthesis of oligomers by stable free radical

polymerization of acrylates, methacrylates, and

styrene with alkoxyamine

initiators

AUTHOR(S): Keul, Helmut; Achten, Dirk; Reining, Birte;

Hocker, Hartwig

CORPORATE SOURCE: Lehrstuhl fur Textilchemie und Makromolekulare

Chemie der Rheinisch-Westfalischen Technischen

Hochschule Aachen, Aachen, 52056, Germany

SOURCE: ACS Symposium Series (2000), 768(Controlled/Living

Radical Polymerization), 408-426 CODEN: ACSMC8; ISSN: 0097-6156

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 03 Sep 2000

AB A controlled polymerization of styrene (St) and Me acrylate (MA) yielding oligomers of Mn ≤ 2000 was achieved using an alkoxyamine as the initiator. For the polymerization of Me methacrylate (MMA) side reactions leading to unsatd. end groups prohibit the control of the polymerization The extent of the side reactions is explained by a sterically hindered combination of TEMPO with the active PMMA chain end. Copolymn. of MA with St reveals an increase of the polymerization rate compared to those of the homopolymns. and a good control of the MA/St copolymer composition within a wide range. The copolymn. parameters were found to be in good agreement with those observed in the free

radical polymerization For the copolymn. of MMA with St a controlled polymerization was achieved at molar fractions of styrene higher than 50 %.

IT 115191-52-1

(synthesis of oligomers by stable free radical polymerization of acrylates, methacrylates, and styrene with alkoxyamine initiators)

RN 115191-52-1 HCAPLUS

CN Propanoic acid, 2-methyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]-, methyl ester (CA INDEX NAME)

CC 35-3 (Chemistry of Synthetic High Polymers)

IT Polymerization

Polymerization catalysts

(radical; synthesis of oligomers by stable free radical polymerization of acrylates, methacrylates, and styrene with alkozyamine initiators)

IT Polydispersity

(synthesis of oligomers by stable free radical polymerization of acrylates, methacrylates, and styrene with alkoxyamine initiators)

IT 2564-83-2, TEMPO 115191-52-1 154554-67-3 212128-87-5

(synthesis of oligomers by stable free radical polymerization of acrylates, methacrylates, and styrene with alkozyamine initiators)

JT 9003-21-8P, Poly(methyl acrylate) 9003-53-6P, Polystyrene 9011-14-7P, PMMA 25034-86-0P, Methyl methacrylate-styrene copolymer 25036-19-5P, Methyl acrylate-styrene copolymer

(synthesis of oligomers by stable free radical polymerization of acrylates, methacrylates, and styrene with alkoxyamine initiators)

REFERENCE COUNT:

CORPORATE SOURCE:

55 THERE ARE 55 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 30 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2000:417358 HCAPLUS Full-text DOCUMENT NUMBER: 133:164347

TITLE: Rate Constants for the Trapping of Various Carbon-Centered Radicals by Nitroxides:

Unimolecular Initiators for Living Free Radical

Polymerization

AUTHOR(S): Skene, W. G.; Scaiano, J. C.; Listigovers, Nancy

A.; Kazmaier, Peter M.; Georges, Michael K. Department of Chemistry, University of Ottawa,

Ottawa, ON, K1N 6N5, Can.

SOURCE: Macromolecules (2000), 33(14), 5065-5072

CODEN: MAMOBX; ISSN: 0024-9297

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 23 Jun 2000

AB The recapping kinetics of a series of unimol. initiators for free radical polymerization were investigated. The rate consts. for the trapping of various carbon-centered radicals were analyzed as a function of structurally different nitroxides in order to determine suitable candidates for living free radical polymerization (LFRP). The technique of laser flash photolysis was used to determine the trapping rate consts. for the radicals in the presence of various amts. of nitroxide and to also gain insight into the properties of the LFRP process.

IT 81913-53-3P

(rate consts. for trapping of carbon-centered radicals by nitroxides and living radical polymerization)

RN 81913-53-3 HCAPLUS

CN Benzeneethanol,  $\beta$ -[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]-, 1-benzoate (CA INDEX NAME)

CC 35-2 (Chemistry of Synthetic High Polymers)

IT 81913-53-3P 117174-68-2P 154554-67-3P 183194-54-9P 212128-87-5P 288098-79-3P

(rate consts. for trapping of carbon-centered radicals by

nitroxides and living radical polymerization)

REFERENCE COUNT: 69 THERE ARE 69 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L68 ANSWER 31 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2000:244986 HCAPLUS Full-text

DOCUMENT NUMBER: 133:4352

TITLE: Tin-free radical cyclization reactions using the

persistent radical effect

AUTHOR(S): Studer, Armido

CORPORATE SOURCE: Lab. Org. Chem., Eidg. Tech. Hochsch., Zurich,

8092, Switz.

SOURCE: Angewandte Chemie, International Edition (2000),

39(6), 1108-1111

CODEN: ACIEF5; ISSN: 1433-7851

PUBLISHER: Wiley-VCH Verlag GmbH

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 133:4352

ED Entered STN: 17 Apr 2000

GΙ

The persistent radical effect (the general principle that explains the highly specific formation of cross-coupling product R1-R2 between two radicals R1 and R2 when one species is long-lived and the other is transient and they are formed at equal rates) was exhibited in the isomerization of I to IV (up to 70% yield) in t-BuOH in presence of camphorsulfonic acid. Ionic and solvent-cage mechanisms were excluded. Scope and limitations of the reaction were examined by introducing Ph substituents and by varying the Ph group (e.g., 2-thienyl, 2-pyridyl) and the nitroxide moiety. Addnl. demonstrations of the superiority of the present alkoxyamine-mediated cyclization over the Bu3SnH-mediated reaction were presented.

IT 270901-39-8P

(isomerization; tin-free radical cyclization reactions following C-O bond homolysis of (alkenyloxy)amines derived from long-lived nitroxides as a demonstration of the persistent radical effect)

RN 270901-39-8 HCAPLUS

CN Piperidine, 2,2,6,6-tetramethyl-1-[(1-phenyl-5-hexen-1-yl)oxy]- (CA INDEX NAME)

Me Me Ph 
$$CH_2$$
  $3-CH$   $CH_2$ 

CC 22-5 (Physical Organic Chemistry)

IT 270901-39-8P 270901-51-4P 270901-52-5P

270901-53-6P 270901-54-7P 270901-55-8P 270901-56-9P 270901-79-6P 270901-83-2P 270901-84-3P 270901-85-4P

270901-86-5P

(isomerization; tin-free radical cyclization reactions following C-O bond homolysis of (alkenyloxy)amines derived from long-lived nitroxides as a demonstration of the persistent radical effect)

IT 270902-07-3P

(tin-free radical cyclization reactions following C-O bond homolysis of (alkenyloxy) amines derived from long-lived nitroxides as a demonstration of the persistent radical effect)

REFERENCE COUNT:

THERE ARE 47 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 32 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2000:215991 HCAPLUS Full-text DOCUMENT NUMBER: 132:250988

47

TITLE: One-step preparation of vinyl group-containing

alkoxyamines as radical generators Hayashi, Masaki; Nakamura, Tomoyuki

INVENTOR(S): Hayashi, Masaki; Nakamura, Tomoyuki PATENT ASSIGNEE(S): Nippon Oil and Fats Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 12 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PA	PATENT NO.								APPLICATION NO.						DATE		
	2000 2000	0957	44		А											9990623 0000517	
,,,		AE, CR, HR, LT, RU,	AG, CU, HU, LU, SD,	AL, CZ, ID, LV, SE,	AM, DE, IL, MA,	AT, DK, IN, MD, SI,	AU, DM, IS, MG, SK,	AZ, DZ, KE, MK,	BA, EE, KG, MN,	BB, ES, KP, MW,	BG, FI, KR, MX,	BR, GB, KZ, NO,	BY, GD, LC, NZ,	CA, GE, LK, PL,	CH, GH, LR, PT,	CN, GM, LS, RO,	
	R₩:	GH, DE,	GM, DK,	KE, ES,	LS, FI,	MW, FR,	SD, GB, GA,	GR,	IE,	IT,	LU,	MC,	NL,	PT,	SE,	•	
AU	2000	0461	42	·	A	ĺ	2001	0109		AU 2	000-	4614.	2	·	2	0000517	
	1188															0000517	
EP	1188	742			В1		2004	1215									
	R:		,	,	•	,	ES, FI,	•	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	
AT	2848									AT 2	000-	9277	96		2	0000517	
	6919															0000517	
PRIORIT	Y APP	LN.	INFO	.:						JP 1	998-	2061	46		A 1	9980722	
										JP 1	999-	1774	22		A 1	9990623	
										JP 1	999-	2420	52		A 1	9990827	
									,	WO 2	000-	JP31	70	,	W 2	0000517	

OTHER SOURCE(S): CASREACT 132:250988; MARPAT 132:250988

ED Entered STN: 04 Apr 2000

AB R1CO2CH2CH(ONR3R4)C6H4CH:CH2 [R1 = aryl; R2O; R2 = C1-12 linear or branched alkyl, cycloalkyl; R3, R4 = C4-6 tertiary alkyl; NR3R4 may form (un)substituted C8-16 ring], useful as polymerization initiators, antioxidants, and scorch inhibitors for polymers (no data), are prepared by heating divinylbenzene with nitroxides and organic peroxides to the thermal decomposition temperature of the peroxides. The products are stable in ionic reactions (no data). OHCH2CH(ONR3R4)C6H4CH:CH2 (R3, R4 = same as above), useful in functional group introduction or as ring-opening polymerization initiators (no data) are prepared by hydrolysis of the above products. 2,2,6,6-Tetramethylpiperidine-1-oxyl was treated with 1,4-divinylbenzene and Bz2O2 at 95° for 3.5 h to give 42.3% PhCO2CH2CH(OZ)C6H4CH:CH2-4 (Z = 2,2,6,6-tetramethylpiperidino).

IT 262616-03-5P

(one-step preparation of vinyl group-containing alkoxyamines as radical generators)

RN 262616-03-5 HCAPLUS

CN Benzeneethanol, 4-ethenyl- $\beta$ -[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]-, benzoate (ester) (9CI) (CA INDEX NAME)

IC ICM C07C239-20

ICS C07D211-94; C08F004-32

CC 25-18 (Benzene, Its Derivatives, and Condensed Benzenoid Compounds)

Section cross-reference(s): 35, 39, 67

IT 262616-03-5P 262616-04-6P 262616-05-7P

262616-06-8P 262616-07-9P 262616-08-0P

262616-09-1P 262616-10-4P 262616-11-5P

262616-12-6P 262846-44-6P 262846-45-7P 262846-46-8P

(one-step preparation of vinyl group-containing alkoxyamines as radical generators)

L68 ANSWER 33 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2000:208068 HCAPLUS Full-text

DOCUMENT NUMBER: 132:334849

TITLE: A "penultimate" initiator for living

free radical polymerization

AUTHOR(S): Skene, W. G.; Scaiano, J. C.; Yap, Glenn P. A.

CORPORATE SOURCE: Department of Chemistry, University of Ottawa,

Ottawa, ON, K1N 6N5, Can.

SOURCE: Polymer Preprints (American Chemical Society,

Division of Polymer Chemistry) (2000), 41(1),

119-120

CODEN: ACPPAY; ISSN: 0032-3934

PUBLISHER: American Chemical Society, Division of Polymer

Chemistry

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 31 Mar 2000

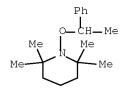
AB To prevent disproportionation reaction of common TEMPO-based initiators (TEMPO = tetramethylpiperidine N-oxide) in living free radical polymerization, a new styrene dimer-connected TEMPO initiator was synthesized. The origins of disproportionation in addition to the kinetic and thermodn. properties were examined The 1-(2,2,6,6-tetramethylpiperidine-N-oxide)-1,3-diphenylpropane initiator was studied by absorption spectroscopy after irradiation, determination of rate constant for radical trapping, and activation parameters of thermal decomposition

IT 154554-67-3P

(living free radical polymerization kinetics in presence of TEMPO-based initiators)

RN 154554-67-3 HCAPLUS

CN Piperidine, 2,2,6,6-tetramethyl-1-(1-phenylethoxy)- (CA INDEX NAME)



CC 35-3 (Chemistry of Synthetic High Polymers)

ST radical polymn modified TEMPO initiator; TEMPO diphenylpropane prepn polymn initiator decompn kinetics trapping

IT Reaction kinetics

(homolytic; living free radical polymerization kinetics in presence of TEMPO-based initiators)

IT Trapping

(kinetics; living free radical polymerization kinetics in presence of TEMPO-based initiators)

IT Decomposition kinetics

Rotational energy

(living free radical polymerization kinetics in presence of TEMPO-based initiators)

IT Polymerization catalysts

(radical; living free radical polymerization kinetics in presence of TEMPO-based initiators)

IT 72379-98-7, 1,3-Diphenylpropyl

(living free radical polymerization kinetics in presence of TEMPO-based initiators)

IT 154554-67-3P

(living free radical polymerization kinetics in presence of TEMPO-based initiators)

IT 124251-79-2P

(preparation of TEMPO-based initiators TEMPO-based initiators for living free radical polymerization)

IT 2564-83-2, TEMPO

(preparation of TEMPO-based initiators TEMPO-based initiators for living free radical polymerization)

IT 1081-75-0, 1,3-Diphenylpropane

(preparation of TEMPO-based initiators TEMPO-based initiators for living free radical polymerization)

REFERENCE COUNT:

11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 34 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1999:757979 HCAPLUS <u>Full-text</u>

DOCUMENT NUMBER: 132:108432

TITLE: High-Yield Synthesis of Functionalized

Alkoxyamine Initiators and

Approach to Well-Controlled Block Copolymers Using

Them

AUTHOR(S): Miura, Yozo; Hirota, Kenichi; Moto, Hiroaki;

Yamada, Bunichiro

CORPORATE SOURCE: Department of Applied Chemistry Faculty of

Engineering, Osaka City University, Sumiyoshi-ku

Osaka, 558-8585, Japan

SOURCE: Macromolecules (1999), 32(25), 8356-8362

CODEN: MAMOBX; ISSN: 0024-9297

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 01 Dec 1999

AΒ Reaction of ethylbenzene, 4-bromo-, 4-(ethoxycarbonyl)-, and 4methoxyethylbenzenes with di-tert-Bu diperoxyoxalate at 35 °C in the presence of stable nitroxide radicals gave alkoxyamines in 50-91% yields. Lithiation of the 4-bromophenylalkoxyamine and subsequent treatment with DMF gave 4formylphenylalkoxyamine in 95% yield, and the reduction of the 4formylphenylalkoxyamine with NaBH4 yielded 4-(hydroxymethyl)phenylalkoxyamine in 89% yield. Anionic polymerization of butadiene (BD) with sec-BuLi and subsequent termination with 4-formylphenylalkoxyamine gave an endfunctionalized poly(butadiene) [poly(BD)]. "Living" radical polymerization of styrene (St) initiated by the functionalized poly(BD) at 120 °C gave a poly(BD)-block-poly(St) with a Mw/Mn of 1.30. Anionic polymerization of hexamethylcyclotrisiloxane (D3) with 4-lithiophenylalkoxyamine gave poly(D3) with an alkoxyamine moiety at a polymer end. "Living" radical polymerization of St initiated by the functionalized poly(D3) gave poly(D3)-block-poly(St) with Mw/Mn's of 1.73-1.80.

IT 154554-67-3P

(high-yield synthesis of functionalized alkozyamine initiators and their use in block polymerization)

RN 154554-67-3 HCAPLUS

CN Piperidine, 2,2,6,6-tetramethyl-1-(1-phenylethoxy)- (CA INDEX NAME)

CC 35-8 (Chemistry of Synthetic High Polymers)

IT Amines, preparation

(alkoxy-; high-yield synthesis of functionalized alkoxyamine initiators and their use in block polymerization)

IT Polysiloxanes, preparation

(block copolymers with styrene; high-yield synthesis of functionalized alkoxyamine initiators and their use in block polymerization)

IT Polymerization catalysts

(high-yield synthesis of functionalized alkoxyamine initiators and their use in block polymerization)

IT 154554-67-3P 178625-97-3P 183194-54-9P 209550-19-6P 209550-20-9P 209550-21-0P 209550-24-3P 209550-25-4P

(high-yield synthesis of functionalized alkoxyamine initiators and their use in block polymerization)

IT 4748-78-1

(high-yield synthesis of functionalized alkozyamine initiators and their use in block polymerization)

IT 209550-22-1P 209550-23-2P

(high-yield synthesis of functionalized alkoxyamine

initiators and their use in block polymerization)

IT 106107-54-4P, 1,3-Butadiene-styrene block copolymer 109668-88-4P,

Hexamethylcyclotrisiloxane-styrene block copolymer

(high-yield synthesis of functionalized alkoxyamine initiators and their use in block polymerization)

IT 9003-17-2DP, Polybutadiene, alkoxyamine-terminated 9003-53-6P, Polystyrene

(in block polymer preparation; high-yield synthesis of functionalized alkoxyamine initiators and their use in block polymerization)

IT 25084-99-5DP, Hexamethylcyclotrisiloxane homopolymer,

alkoxyamine-terminated

(in block polymer preparation; high-yield synthesis of functionalized alkoxyamine initiators and their use in block polymerization)

IT 100-41-4, Ethylbenzene, reactions 1876-22-8 2564-83-2, TEMPO (in catalyst preparation; high-yield synthesis of functionalized alkoxyamine initiators and their use in block polymerization)

REFERENCE COUNT: 34 THERE ARE 34 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 35 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1999:595167 HCAPLUS Full-text

DOCUMENT NUMBER: 131:229163

TITLE: Nitroxyl derivatives with glycidyl or

alkylcarbonyl groups as initiators for radical

polymerization

INVENTOR(S): Fuso, Francesco; Roth, Michael; Wunderlich, Wiebke

PATENT ASSIGNEE(S): Ciba Specialty Chemicals Holding Inc., Switz.

SOURCE: PCT Int. Appl., 36 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PA:	TENT	NO.			KIND DATE		APPLICATION NO.						DATE			
WO	9946	 261			A1		 1999	0916							1	9990226
	W:	AL,	AM,	ΑT,	ΑU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	CA,	CH,	CN,	CU,	CZ,
		DE,	DK,	EE,	ES,	FΙ,	GB,	GD,	GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,
		IS,	JP,	KE,	KG,	KP,	KR,	KΖ,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	MD,
		MG,	MK,	MN,	MW,	MX,	NO,	NZ,	PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,
		SK,	SL,	ТJ,	TM,	TR,	TT,	UA,	UG,	US,	UZ,	VN,	YU,	ZW		
	RW:	GH,	GM,	KE,	LS,	MW,	SD,	SL,	SZ,	UG,	ZW,	AT,	BE,	CH,	CY,	DE,
		DK,	ES,	FI,	FR,	GB,	GR,	IE,	IT,	LU,	MC,	NL,	PT,	SE,	BF,	BJ,
		CF,	CG,	CI,	CM,	GA,	GN,	GW,	ML,	MR,	ΝE,	SN,	TD,	ΤG		
TW	4955	15			В		2002	0721		TW 1	999-	8810	0735		1	9990118
CA	2321	792			A1		1999	0916	1	CA 1	999-	2321	792		1	9990226
ΑU	9929	296			Α		1999	0927		AU 1	999-	2929	6		1	9990226
EP	1071	681			A1		2001	0131		EP 1	999-	9102	86		1	9990226
EP	1071	681			В1		2006	0830								
	R:	BE,	DE,	ES,	FR,	GB,	ΙΤ,	NL								
JP	2002	5060	70		Τ		2002	0226	1	JP 2	000 -	5356	40		1	9990226
IN	1999	MA00.	275		Α		2007	0511		IN 1	999-1	MA27.	5		1	9990308
US	6566	468			В1		2003	0520		US 2	000-	6235	47		2	0000905
MX	X 2000PA08767 A 20010328 MX						MX 2	000 - 1	PA87	67		2	0000908			
US 2002107397 A1 20020808 US								3 US 2001–13884 20011211								

US 6683142	В2	20040127				
US 2003220423	A1	20031127	US	2003-422497		20030424
US 6927295	В2	20050809				
US 2005131177	A1	20050616	US	2005-35915		20050114
PRIORITY APPLN. INFO.:			EP	1998-810194	А	19980309
			ΕP	1998-810531	А	19980611
			WO	1999-EP1233	W	19990226
				1000 060001	- 0	10000001
			US	1999-262804	А3	19990304
				0001 10004	7.0	00011011
			US	2001-13884	АЗ	20011211
			110	2002 422407	71 7	20020424
			US	2003-422497	АЗ	20030424

OTHER SOURCE(S): MARPAT 131:229163

ED Entered STN: 21 Sep 1999

AB Cyclic and linear nitroxyl derivs. having glycidyl or alkylcarbonyl groups attached to the O of the nitroxyl group such as are useful as initiators for radical polymerization of ethylenically unsatd. monomers to give polymers with narrow polydispersity at high monomer conversion, relatively low temps., and short reaction times. A typical initiator was manufactured by adding a solution containing 50 g 2-(4- ethylphenoxyethyl)oxirane (I) and 26.4 g tert-Bu hydroperoxide in 30 min to a mixture containing I 57, 4-propoxy-2,2,6,6-tetramethylpiperidine 1-oxyl 10.7, and MoO3 0.72 g at 70° with stirring, reducing the pressure to 200 millibar, and heating 18 h at 100°.

IT 243972-01-2P

IC

(nitroxyl derivs. with glycidyl or alkylcarbonyl groups as initiators for radical polymerization)

RN 243972-01-2 HCAPLUS

ICM C07D405-12

CN Piperidine, 2,2,6,6-tetramethyl-1-[1-[4-(oxiranylmethoxy)phenyl]ethoxy ]-4-propoxy- (9CI) (CA INDEX NAME)

ICS C07D211-94; C08F004-00
CC 35-3 (Chemistry of Synthetic High Polymers)
 Section cross-reference(s): 23, 27
IT 243972-01-2P 243972-02-3P 243972-03-4P

IT 243972-01-2P 243972-02-3P 243972-03-4P 243972-04-5P 243972-05-6P 243972-06-7P 243972-07-8P 243972-08-9P 243972-10-3P 243972-11-4P 243972-12-5P 243972-13-6P 243972-14-7P 243972-15-8P 243972-16-9P 243972-17-0P 243972-18-1P

(nitroxyl derivs. with glycidyl or alkylcarbonyl groups as initiators for radical polymerization)

REFERENCE COUNT: 15 THERE ARE 15 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 36 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1999:317797 HCAPLUS Full-text

DOCUMENT NUMBER: 131:102577

TITLE: Synthesis and characterization of  $$\omega$-unsaturated poly(styrene-b-n-butyl$ 

methacrylate) block copolymers using
TEMPO-mediated controlled radical polymerization
Burguiere, Carine; Dourges, Marie-Anne; Charleux,

Dangalette, Vainer Jaco Diame

Bernadette; Vairon, Jean-Pierre

CORPORATE SOURCE: Laboratoire de Chimie Macromoleculaire, Unite

Mixte associee au CNRS UMR 7610, Universite Pierre

et Marie Curie, Paris, 75252, Fr.

SOURCE: Macromolecules (1999), 32(12), 3883-3890

CODEN: MAMOBX; ISSN: 0024-9297

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 25 May 1999

AUTHOR(S):

AB Bu methacrylate has been polymerized in bulk at  $130^{\circ}$ C in the presence of given amts. of a nitroxide stable free radical (TEMPO = 2, 2, 6, 6-

tetramethylpiperidine-N-oxyl) using either a low mol. weight alkozyamine initiator [2,2,6,6-tetramethyl-1-(1- phenethyloxy)piperidine] or a TEMPOcapped polystyrene macroinitiator. Complete consumption of both initiators was always observed In contrast, very low final monomer conversions were found. Proton NMR spectroscopy and MALDI-TOF mass spectrometry were used for investigation of the polymer structure. They showed that the formed poly(Bu methacrylate) had the attached initiator at one end and that a block copolymer was synthesized when the polystyrene macroinitiator was used. The other terminal functionality of the polymer was not a TEMPO-based alkoxyamine but a methylene unsatn. exclusively. Particularly, no saturated polymer which would also be formed by conventional disproportionation reaction between two propagating radicals could be detected. From this result, it was concluded that the main chain-breaking event is the  $\beta$ -hydrogen transfer from a propagating radical to TEMPO (also called disproportionation reaction). initial concentration of added TEMPO was shown to directly influence the poly(Bu methacrylate) block length independently of the initial concentration of alkoxyamine: the larger the concentration of TEMPO, the shorter the block length. The rate constant of disproportionation was calculated to be kdis = 1.4 + 106 L mol - 1 s - 1 at 130 °C.

IT 154554-67-3

(diblock polymer preparation using TEMPO-mediated controlled radical polymerization)  $\,$ 

RN 154554-67-3 HCAPLUS

CN Piperidine, 2,2,6,6-tetramethyl-1-(1-phenylethoxy)- (CA INDEX NAME)

CC 35-3 (Chemistry of Synthetic High Polymers)

IT 2564-83-2 154554-67-3

(diblock polymer preparation using TEMPO-mediated controlled radical

polymerization)

REFERENCE COUNT: 32 THERE ARE 32 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L68 ANSWER 37 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 1999:211299 HCAPLUS Full-text

DOCUMENT NUMBER: 130:352729

TITLE: Novel methods for the fabrication of well-defined

and patterned polymer brushes

AUTHOR(S): Benoit, Didier; Husemann, Marc; Mecerreyes, David;

Morrison, Michael; Hinsberg, William; Hawker, Craig J.; Hedrick, James L.; Shah, Rahul; Abbott,

Nicholas L.

CORPORATE SOURCE: IBM Almaden Research Center, San Jose, CA,

95120-6099, USA

SOURCE: Polymer Preprints (American Chemical Society,

Division of Polymer Chemistry) (1999), 40(1),

498-499

CODEN: ACPPAY; ISSN: 0032-3934

PUBLISHER: American Chemical Society, Division of Polymer

Chemistry

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 05 Apr 1999

Strategies were developed for the preparation of patterned polymer brushes, AΒ either isolated structures from microcontact printed Au surfaces, or chemical patterning of continuous brush structures using lithog. techniques. A nonreactive self-assembled monolayer [SAM] of CH3(CH2)15SH is microcontact printed onto a gold surface, then HO(CH2CH2O)2(CH2)11SH was selectively assembled onto the bare regions of the gold surface. The final step is surface initiated ring opening polymerization of  $\epsilon$ -caprolactone or polymerization of other monomers from the functionalized areas of the patterned SAM based on the hydroxyl groups as initiators. Another strategy involves the patterning of a continuous polymer brush into areas of hydrophilic and hydrophobic chains; an alkoxyamine initiator can be readily hydrosilylated with trichlorosilane to obtain the trichlorosiliyl derivative This derivative can be attached to a variety of surfaces, e.g., native silicon oxide layer of silicon wafers by reaction with the surface silanol groups. Polymer brushes can then be grown on the functionalized surface from vinyl monomers. Patterning of polymer brushes involved spin coating a photoresist layer on the polymer brush, exposure of the photoresist to deep-UV irradiation through a mask, forming a photoacid that diffuses to the polymer brush. Removal of the photoresist led to the patterned brush.

IT 225229-07-2P

(novel methods for fabrication of well-defined and patterned polymer brushes)

RN 225229-07-2 HCAPLUS

CN Piperidine, 2,2,6,6-tetramethyl-1-[1-[4-[[6-

(trichlorosily1)hexy1]oxy]methy1]pheny1]ethoxy]- (CA INDEX NAME)

CC 35-7 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 36

225229-07-2P ΙT

> (novel methods for fabrication of well-defined and patterned polymer brushes)

821-41-0, Hex-5-en-1-ol 10025-78-2, Trichlorosilane TT 212132-38-2

> (novel methods for fabrication of well-defined and patterned polymer brushes)

225229-06-1P ΙT

> (novel methods for fabrication of well-defined and patterned polymer brushes)

REFERENCE COUNT:

THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 38 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN 1999:211107 HCAPLUS Full-text ACCESSION NUMBER:

DOCUMENT NUMBER: 130:352602

TITLE: Preparation of alkozyamine

19

initiators carrying a functional group

Miura, Yozo; Hirota, Kenichi; Moto, Hiroaki; AUTHOR(S):

Yamada, Bunichiro

CORPORATE SOURCE: Department of Applied Chemistry, Faculty of

Engineering, Osaka City University, Osaka,

558-8585, Japan

SOURCE: Polymer Preprints (American Chemical Society,

Division of Polymer Chemistry) (1999), 40(1),

123-124

CODEN: ACPPAY; ISSN: 0032-3934

PUBLISHER: American Chemical Society, Division of Polymer

Chemistry

DOCUMENT TYPE: Journal LANGUAGE: English

ΕD Entered STN: 05 Apr 1999

AΒ Alkoxyamines useful as catalysts for living radical polymerization are prepared in high yields by the reaction of ethylbenzene with di-tert-Bu peroxalate at 35° in the presence of stable nitroxides such as TEMPO and ditert-Bu nitroxide. By using this procedure functionalized alkoxyamines including p-bromo substituted alkoxyamine were also prepared in high yield. Furthermore, p-Formyl and p-hydroxymethyl substituted alkoxyamines were obtained from the p-bromo substituted alkoxyamine.

ΙT 178625-97-3P

(catalysts; preparation of alkoxyamine initiators

for living radical polymerization)

RN 178625-97-3 HCAPLUS

CN Piperidine, 1-[1-(4-bromophenyl)ethoxy]-2,2,6,6-tetramethyl- (CA INDEX NAME)

```
CC
     35-3 (Chemistry of Synthetic High Polymers)
ΙT
     Polymerization catalysts
     Polymerization catalysts
        (living, radical; preparation of alkoxyamine
        initiators containing functional group)
ΙT
     178625-97-3P 209550-20-9P 209550-21-0P
     209550-22-1P 209550-23-2P 209550-24-3P
        (catalysts; preparation of alkoxyamine initiators
        for living radical polymerization)
     68-12-2, DMF, reactions
                              100-41-4, Ethylbenzene, reactions
ΤТ
     594-19-4, tert-Butyllithium
                                   1876-22-8, Di-tert-butyl peroxalate
     2406-25-9, Di-tert-butyl nitroxide
                                         2564-83-2, TEMPO
     2,2,5,5-Tetramethylpyrrolidinyl-1-oxyl
        (in preparation of alkoxyamine initiators for living
        radical polymerization)
ΙT
     9003-53-6P, Polystyrene
        (preparation of in presence of alkoxyamine initiators
REFERENCE COUNT:
                         17
                               THERE ARE 17 CITED REFERENCES AVAILABLE FOR
                               THIS RECORD. ALL CITATIONS AVAILABLE IN THE
                               RE FORMAT
L68 ANSWER 39 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:
                         1999:87261 HCAPLUS Full-text
DOCUMENT NUMBER:
                         130:237921
                         Direct Synthesis of Dispersed Nanocomposites by in
TITLE:
                         Situ Living Free Radical Polymerization Using a
                         Silicate-Anchored Initiator
AUTHOR(S):
                         Weimer, Marc W.; Chen, Hua; Giannelis, Emmanuel
                         P.; Sogah, Dotsevi Y.
CORPORATE SOURCE:
                         Department of Chemistry and Chemical Biology Baker
                         Laboratory Department of Materials Science and
                         Engineering, Cornell University, Ithaca, NY,
                         14853, USA
                         Journal of the American Chemical Society (1999),
SOURCE:
                         121(7), 1615-1616
                         CODEN: JACSAT; ISSN: 0002-7863
                         American Chemical Society
PUBLISHER:
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
     Entered STN: 11 Feb 1999
ED
     Anchoring a living free radical polymerization (LFRP) initiator inside the
AB
     galleries of layered silicate hosts followed by intercalation and
     polymerization of styrene gives directly dispersed polystyrene (PS) -silicate
     nanocomposite. The initiator was prepared and ion-exchanged onto a com.
     montmorillonite layered silicate to obtain the intercalated species. The LFRP
     was carried out by heating a dispersion of the intercalated initiator species
     in styrene for 4 h; the system solidified completely to yield the
     nanocomposite of silicate randomly dispersed spatially and directionally in
     the polystyrene matrix consisting of small domains. This level of uniform
     dispersion is not achievable by either melt or solution intercalation of a
     preformed polystyrene. The polymer was desorbed from the silicate by
     refluxing the nanocomposite in THF/LiBr; the low polydispersity index (PDI) of
     1.3 and the agreement between the calculated number-average mol. weight (Mn)
     of 24 400 and observed Mn of 21 500, indicate a remarkably well-behaved
     reaction even under such heterogeneous conditions.
     221362-46-5P
ΤT
        (radical initiator; preparation of radical initiator
```

for synthesis of dispersed silicate-polystyrene nanocomposites)

RN 221362-46-5 HCAPLUS

CN Benzenemethanaminium, 4-[2-(benzoyloxy)-1-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl]-N,N,N-trimethyl-, chloride (9CI) (CA INDEX NAME)

● C1-

CC 35-4 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 37, 57

ST polystyrene layered silicate hybrid nanocomposite anchoring initiator

IT Polymerization

Polymerization catalysts

(radical; synthesis of dispersed nanocomposites by in situ living free radical polymerization using silicate-anchored initiator)

IT Dispersion (of materials)

Hybrid organic-inorganic materials

Intercalation

Ion exchange

Polydispersity

(synthesis of dispersed nanocomposites by in situ living free radical polymerization using silicate-anchored initiator)

IT 9003-53-6P

(nanocomposites; synthesis of dispersed nanocomposites by in situ living free radical polymerization using silicate-anchored initiator)

IT 75-50-3, reactions

(preparation of radical initiator for synthesis of dispersed silicate-polystyrene nanocomposites)

IT 221362-46-5P

(radical initiator; preparation of radical initiator

for synthesis of dispersed silicate-polystyrene nanocomposites)

IT 216104-33-5

(radical initiator; preparation of radical initiator

for synthesis of dispersed silicate-polystyrene nanocomposites)

IT 1318-93-0, Montmorillonite, uses

(support and nanocomposite component; synthesis of dispersed nanocomposites by in situ living free radical polymerization using silicate-anchored initiator)

REFERENCE COUNT:

THERE ARE 32 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 40 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1999:76038 HCAPLUS Full-text DOCUMENT NUMBER: 130:237911

32

TITLE: Controlled Synthesis of Polymer Brushes by

"Living" Free Radical Polymerization Techniques Husseman, Marc; Malmstroem, Eva E.; McNamara, Molly; Mate, Mathew; Mecerreyes, David; Benoit,

Didier G.; Hedrick, James L.; Mansky, Paul; Huang,

E.; Russell, Thomas P.; Hawker, Craig J.

CORPORATE SOURCE: Center for Polymeric Interfaces and Macromolecular

Assemblies, IBM Almaden Research Center, San Jose,

CA, 95120-6099, USA

SOURCE: Macromolecules (1999), 32(5), 1424-1431

CODEN: MAMOBX; ISSN: 0024-9297

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 05 Feb 1999

AB The preparation of a wide variety of unique polymer brush structures can be accomplished by living free radical polymerization of vinyl monomers from surface-tethered alkoxyamines or from tethered  $\alpha$ -halo esters in the presence of (PPh3)2NiBr2. The use of a living free radical process permits the mol. weight and polydispersity of the covalently attached polymer chains to be accurately controlled while also allowing the formation of block copolymers by the sequential growth of monomers from the surface. These block and random copolymer brushes were used to control surface properties.

IT 212132-49-5P

AUTHOR(S):

(alkoxyamine precursor; controlled synthesis of polystyrene and polymethacrylate brushes by living free radical polymerization using surface-tethered alkoxyamine initiators)

RN 212132-49-5 HCAPLUS

CN Piperidine, 2,2,6,6-tetramethyl-1-[1-[4-[(4-pentenyloxy)methyl]phenyl]ethoxy]- (9CI) (CA INDEX NAME)

- CC 35-3 (Chemistry of Synthetic High Polymers)
- IT Polymers, preparation

(brush; controlled synthesis of polystyrene and polymethacrylate brushes by living free radical polymerization using surface-tethered alkoxyamine initiators)

IT Polymer chains

(brushes; controlled synthesis of polystyrene and polymethacrylate brushes by living free radical polymerization using surface-tethered alkoxyamine initiators)

IT Hydrosilylation

Polydispersity

(controlled synthesis of polystyrene and polymethacrylate brushes by living free radical polymerization using surface-tethered alkoxyamine initiators)

IT Polymerization

Polymerization catalysts

(radical; controlled synthesis of polystyrene and polymethacrylate brushes by living free radical polymerization using surface-tethered

alkoxyamine initiators)

IT 212132-49-5P 221318-43-0P

(alkoxyamine precursor; controlled synthesis of polystyrene and polymethacrylate brushes by living free radical polymerization using surface-tethered alkoxyamine initiators)

IT 110-05-4, Di-tert-butyl peroxide 16940-66-2, Sodium borohydride 135620-04-1, Jacobsen's catalyst

(controlled synthesis of polystyrene and polymethacrylate brushes by living free radical polymerization using surface-tethered alkoxyamine initiators)

IT 821-09-0, Pent-4-enol 1066-35-9, Dimethylchlorosilane 1592-20-7 2564-83-2, 2,2,6,6-Tetramethylpiperidinyloxy 7646-69-7, Sodium hydride 10025-78-2, Trichlorosilane 212132-38-2

(controlled synthesis of polystyrene and polymethacrylate brushes by living free radical polymerization using surface-tethered alkoxyamine initiators)

IT 9003-53-6P 9011-14-7P, Poly(methyl methacrylate) 26010-51-5P, 2-Hydroxyethyl methacrylate-styrene copolymer 106911-77-7P, Methyl methacrylate-styrene block copolymer

(controlled synthesis of polystyrene and polymethacrylate brushes by living free radical polymerization using surface-tethered alkozyamine initiators)

- IT 600-00-0, Ethyl-2-bromo-2-methylpropionate 154554-67-3 (controlling initiator; controlled synthesis of polystyrene and polymethacrylate brushes by living free radical polymerization using surface-tethered alkoxyamine initiators)
- IT 212132-48-4P 221318-51-0P
   (intermediate; controlled synthesis of polystyrene and
   polymethacrylate brushes by living free radical polymerization using
   surface-tethered alkomyamine initiators)
- IT 14126-37-5

(polymerization catalyst; controlled synthesis of polystyrene and polymethacrylate brushes by living free radical polymerization using surface-tethered alkoxyamine initiators)

IT 821-41-0, 5-Hexen-1-ol 20769-85-1, 2-Bromo-2-methylpropionyl bromide 89359-54-6 161776-41-6, 2-Phenyl-2-[(2,2,6,6-

tetramethylpiperidino)oxy]ethanol

(starting material; controlled synthesis of polystyrene and polymethacrylate brushes by living free radical polymerization using surface-tethered alkoxyamine initiators)

IT 7631-86-9, Silica, reactions

(substrate surface reactive layer; controlled synthesis of polystyrene and polymethacrylate brushes by living free radical polymerization using surface-tethered alkozyamine initiators)

IT 221318-45-2P 221318-47-4P

(surface bound initiator; controlled synthesis of polystyrene and polymethacrylate brushes by living free radical polymerization using surface-tethered alkoxyamine initiators)

IT 221318-53-2P

(surface-bound initiator; controlled synthesis of polystyrene and polymethacrylate brushes by living free radical polymerization using surface-tethered alkoxyamine initiators)

IT 7440-21-3, Silicon, uses

(wafer, substrate; controlled synthesis of polystyrene and polymethacrylate brushes by living free radical polymerization using surface-tethered alkoxyamine initiators)

REFERENCE COUNT: 42 THERE ARE 42 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 41 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1998:509835 HCAPLUS <u>Full-text</u>

DOCUMENT NUMBER: 129:216942

TITLE: Meisenheimer Rearrangement of Allyl N-Oxides as a

Route to Initiators for

Nitroxide-Mediated "Living" Free Radical

Polymerizations

AUTHOR(S): Bergbreiter, David E.; Walchuk, Brian

CORPORATE SOURCE: Department of Chemistry, Texas A&M University,

College Station, TX, 77842, USA

SOURCE: Macromolecules (1998), 31(18), 6380-6382

CODEN: MAMOBX; ISSN: 0024-9297

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 18 Aug 1998

AB A new method was developed for preparing unimol. initiators for the living radical polymerization of styrene based on the oxidation and Meisenheimer rearrangement of allylic N-oxide derivs. of 2,2,6,6-tetramethylpiperidine. The N-allyl derivs. of 2,2,6,6-tetramethylpiperidine were oxidized to the corresponding N-oxide by treatment with m-chloroperbenzoic acid. This product spontaneously rearranges under the reaction conditions to give O-allyl TEMPO derivs. The TEMPO derivs. were evaluated as catalysts for the living radical polymerization of styrene.

IT 102261-92-7P

(polymerization catalyst; Meisenheimer rearrangement in preparation of allyloxy TEMPO derivs. for polymerization catalysts for styrene living radical polymerization)

RN 102261-92-7 HCAPLUS

CN Piperidine, 2,2,6,6-tetramethyl-1-(phenylmethoxy)- (CA INDEX NAME)

CC 35-3 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 27

ST Meisenheimer rearrangement allyloxy TEMPO deriv prepn; polymn catalyst allyloxy TEMPO deriv prepn; styrene living radical polymn catalyst

IT Rearrangement

(Meisenheimer; in preparation of allyloxy TEMPO derivs. for polymerization catalysts for styrene living radical polymerization)

IT Polymerization

(living, radical, kinetics; Meisenheimer rearrangement in preparation of allyloxy TEMPO derivs. for polymerization catalysts for styrene living radical polymerization)

IT Polymerization

Polymerization

(living, radical, mechanism; Meisenheimer rearrangement in preparation of allyloxy TEMPO derivs. for polymerization catalysts for styrene living radical polymerization)

IT Polymerization catalysts
 Polymerization catalysts

Polymerization kinetics (living, radical; Meisenheimer rearrangement in preparation of allyloxy TEMPO derivs. for polymerization datalysts for styrene living radical polymerization) 9003-53-6P, Polystyrene ΙT (Meisenheimer rearrangement in preparation of allyloxy TEMPO derivs. for polymerization catalysts for styrene living radical polymerization) 783-76-6P, 1-Benzyl-2,2,6,6-Tetramethylpiperidine 54262-93-0P, TT 1-Ally1-2,2,6,6-Tetramethylpiperidine 204905-80-6P 204905-84-0P (catalyst intermediate; Meisenheimer rearrangement in preparation of allyloxy TEMPO derivs. for polymerization catalysts for styrene living radical polymerization) 100-39-0, Benzyl bromide 106-95-6, Allyl bromide, reactions ΙT 768-66-1, 2,2,6,6-Tetramethylpiperidine 870-63-3, Prenyl bromide 4392-24-9, Cinnamyl bromide (catalyst starting material; Meisenheimer rearrangement in preparation of allyloxy TEMPO derivs. for polymerization catalysts for styrene living radical polymerization) 100-42-5, processes ΙT (kinetics and polymerization mechanism; Meisenheimer rearrangement in preparation of allyloxy TEMPO derivs. for polymerization catalysts for styrene living radical polymerization) 212576-03-9P ΙT (polymerization catalyst byproduct; Meisenheimer rearrangement in preparation of allyloxy TEMPO derivs. for polymerization datalysts for styrene living radical polymerization) 131531-08-3P 131749-90-1P 102261-92-7P 204906-02-5P TT (polymerization catalyst; Meisenheimer rearrangement in preparation of allyloxy TEMPO derivs. for polymerization catalysts for styrene living radical polymerization) THERE ARE 13 CITED REFERENCES AVAILABLE FOR REFERENCE COUNT: 13 THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L68 ANSWER 42 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN 1998:503834 HCAPLUS Full-text ACCESSION NUMBER: DOCUMENT NUMBER: 129:203301 TITLE: A versatile and efficient synthesis of alkoxyamine LFR initiators via manganese based asymmetric epoxidation catalysts Dao, Julian; Benoit, Didier; Hawker, Craig J. AUTHOR(S): Center for Polymeric Interfaces and Macromolecular CORPORATE SOURCE: Assemblies, IBM Research Division, Almaden Research Center, San Jose, CA, 95120-6099, USA SOURCE: Journal of Polymer Science, Part A: Polymer Chemistry (1998), 36(12), 2161-2167 CODEN: JPACEC; ISSN: 0887-624X John Wiley & Sons, Inc. PUBLISHER: DOCUMENT TYPE: Journal LANGUAGE: English EDEntered STN: 14 Aug 1998 AΒ Styrene derivs, were treated with TEMPO in the presence of manganese complex epoxidn. catalysts to give piperidine ethers that were useful as living free radical polymerization catalysts. ΙT 212132-52-0P (synthesis of alkoxyamine living free radical polymerization initiators via manganese-based asym. epoxidn. catalysts) 212132-52-0 HCAPLUS RN Piperidine, 1-[1-[4-(azidomethyl)phenyl]ethoxy]-2,2,6,6-tetramethyl-CN

(CA INDEX NAME)

CC 35-3 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 27

IT 212132-52-0P

(synthesis of alkoxyamine living free radical polymerization initiators via manganese-based asym. epoxidn. catalysts)

IT 178625-98-4F 184646-30-8F 209550-21-0F 209550-24-3F 212132-38-2F 212132-39-3F 212132-40-6F 212132-41-7F 212132-43-9F 212132-45-1F 212132-47-3F 212132-53-1F

(synthesis of alkoxyamine living free radical polymerization initiators via manganese-based asym. epoxidn. catalysts)

REFERENCE COUNT: 28 THERE ARE 28 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 43 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1998:411135 HCAPLUS <u>Full-text</u>

DOCUMENT NUMBER: 129:95758

TITLE: High-Yield Synthesis of Alkomyamine

Initiators Carrying a Functional Group by Reaction of Ethylbenzenes with Di-tert-butyl Diperoxalate in the Presence of Nitroxides Miura, Yozo; Hirota, Kenichi; Moto, Hiroaki;

Yamada, Bunichiro

CORPORATE SOURCE: Department of Applied Chemistry, Faculty of

Engineering, Osaka City University, Osaka,

558-8585, Japan

SOURCE: Macromolecules (1998), 31(14), 4659-4661

CODEN: MAMOBX; ISSN: 0024-9297

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 08 Jul 1998

AB The reaction of ethylbenzenes with di-tert-Bu diperoxalate in the presence of nitroxides at 35° gives alkoxyamines, including 4-bromo, 4-ethoxycarbonyl, and 4-methoxy derivs. in good yields. Bulk polymerization of styrene initiated by 4-ethoxycarbonyl alkoxyamines was carried out at 120°. The polydispersity of polystyrene obtained is <1.3, indicating that these functionalized alkoxyamines can be used for the syntheses of controlled macromol. architectures.

IT 178625-97-3P

AUTHOR(S):

(preparation of alkozyamine initiators from ethylbenzenes with t-bu-diperoxalate and nitroxides and use in controlled styrene polymerization)

RN 178625-97-3 HCAPLUS

CN Piperidine, 1-[1-(4-bromophenyl)ethoxy]-2,2,6,6-tetramethyl- (CA INDEX NAME)

CC 35-3 (Chemistry of Synthetic High Polymers) Section cross-reference(s): 27, 67

ST alkoxyamine prepn radical polymn initiator; styrene polymn ethoxycarbonyl alkoxyamine initiator

IT Polydispersity

(preparation of alkoxyamine initiators from ethylbenzenes with t-bu-diperoxalate and nitroxides and use in controlled styrene polymerization)

IT Polymerization catalysts

(radical; preparation of alkoxyamine initiators from ethylbenzenes with t-bu-diperoxalate and nitroxides and use in controlled styrene polymerization)

IT 178625-97-3P

(preparation of alkozyamine initiators from ethylbenzenes with t-bu-diperoxalate and nitroxides and use in controlled styrene polymerization)

IT 154554-67-3P, 2,2,6,6-Tetramethyl-1-(1-phenylethoxy)piperidine 209550-19-6P 209550-20-9P 209550-21-0P 209550-24-3P

(preparation of alkoxyamine initiators from ethylbenzenes with t-bu-diperoxalate and nitroxides and use in controlled styrene polymerization)

IT 68-12-2, DMF, reactions 100-41-4, Ethylbenzene, reactions 594-19-4 2406-25-9, DTBN 2564-83-2, TEMPO 3141-58-0, tert-Butoxyl radical 3229-53-6, Proxo 16853-85-3, Lithium aluminum hydride 183194-54-9

(preparation of alkoxyamine initiators from ethylbenzenes with t-bu-diperoxalate and nitroxides and use in controlled styrene polymerization)

IT 209550-22-1P 209550-23-2P

(preparation of alkoxyamine initiators from ethylbenzenes with t-bu-diperoxalate and nitroxides and use in controlled styrene polymerization)

IT 9003-53-6P, Polystyrene 209550-25-4P

(preparation of alkoxyamine initiators from ethylbenzenes with t-bu-diperoxalate and nitroxides and use in controlled styrene polymerization)

REFERENCE COUNT: 17 THERE ARE 17 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L68 ANSWER 44 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1997:142814 HCAPLUS Full-text DOCUMENT NUMBER: 126:186432

TITLE: Reactions of the "stable" nitroxide radical TEMPO.

Relevance to "living" free radical polymerizations

and autopolymerization of styrene

AUTHOR(S): Connolly, Terrence J.; Scaiano, J. C.

CORPORATE SOURCE: Department of Chemistry, University of Ottawa,

Ottawa, ON, K1N 6N5, Can.

SOURCE: Tetrahedron Letters (1997), 38(7), 1133-1136

CODEN: TELEAY; ISSN: 0040-4039

PUBLISHER: Elsevier
DOCUMENT TYPE: Journal
LANGUAGE: English
ED Entered STN: 05 Mar 1997

The stable nitroxide TEMPO reacts with styrene and polystyrene models under conditions typically used for polymns. These results show that the nitroxide is not inert at the elevated temps. necessary to conduct polymns. The implications of H abstraction and olefin addition by the nitroxide are discussed, particularly as they apply to initiation, control of polydispersity

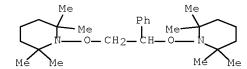
and nitroxide mediated autopolymn. of styrene.

IT 187410-28-2P

(preparation in reaction of styrene with TEMPO)

RN 187410-28-2 HCAPLUS

CN Piperidine, 1,1'-[(1-phenyl-1,2-ethanediyl)bis(oxy)]bis[2,2,6,6-tetramethyl- (CA INDEX NAME)



CC 35-3 (Chemistry of Synthetic High Polymers)

ST TEMPO mediated living polymn styrene; mechanism TEMPO mediated living polymn styrene; initiator TEMPO living polymn styrene

IT Polymerization

Polymerization catalysts Polymerization catalysts

(living, radical; mechanism of TEMPO-mediated living autopolymn. of styrene)

IT 187410-28-2P

(preparation in reaction of styrene with TEMPO)

REFERENCE COUNT: 23 THERE ARE 23 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L68 ANSWER 45 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1991:492290 HCAPLUS Full-text

DOCUMENT NUMBER: 115:92290

TITLE: Preparation of (aminooxycarbonyl)aryloxypyrimidine

s and -arylthiopyrimidines as herbicides

INVENTOR(S): Hiratsuka, Mitsunori; Hirata, Naonori; Saitoh,

Kazuo; Shibata, Hideyuki

PATENT ASSIGNEE(S): Sumitomo Chemical Co., Ltd., Japan

SOURCE: Eur. Pat. Appl., 80 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 426476	A1	19910508	EP 1990-311985	19901101

EP	426476				В1	B1 19940727						
	R:	BE,	CH,	DE,	DK,	ES,	FR,	GB,	IT, L	I, NL		
US	5125	957			А	1	1992	0630	US	1990-602689		19901024
CA	2029	027			A1	1	1991	0502	CA	1990-2029027		19901031
AU	9065	703			А	1	1991	0509	AU	1990-65703		19901031
AU	6299	99			В2	1	1992	1015				
CN	1051	556			Α	1	1991	0522	CN	1990-108773		19901031
CN	1028	714			В	1	L995	0607				
HU	5560	3			A2	1	1991	0628	HU	1990-6966		19901031
HU	2082	30			В	1	L993	0928				
BR	9005	540			Α	1	L991	0917	BR	1990-5540		19901031
RU	2029	471			C1	1	L995	0227	RU	1990-4831578		19901031
JP	0417	8371			Α	1	L992	0625	JP	1990-297558		19901101
ES	2057	437			Т3	1	1994	1016	ES	1990-311985		19901101
US	5300	640			Α	1	1994	0405	US	1991-810166		19911219
AU	9221	139			Α	1	1992	1022	AU	1992-21139		19920820
AU	6445	38			В2	1	1993	1209				
PRIORITY	APP	LN.	INFO	.:					JP	1989-287133	А	19891101
									JP	1990-142390	A	19900530
									JP	1990-213934	А	19900810
									01	1000 210001		10000010
									US	1990-602689	АЗ	19901024

OTHER SOURCE(S): MARPAT 115:92290

ED Entered STN: 06 Sep 1991

GΙ

Title compds. [I; R1, R2 = H, alkyl, alkenyl, alkynyl, haloalkyl, alkoxyalkyl, alkenyloxyalkyl, alkoxycarbonylalkyl, cyanoalkyl, (CH2)nCONR5R6, (CH2)nS(O)mR7, etc.; R3, R4 = alkyl, alkoxy, haloalkoxy, halo; R5, R6 = H, alkyl, alkenyl, alkynyl; R7 = alkyl, alkenyl, alkynyl; m = 0-2; n = 1-4; Y1-Y3 = H, halo, alkyl, alkoxy; X = 0, S; Z = N, CY4; Y4 = H, OH, SH, NO2, halo, alkyl, alkenyl, alkynyl, alkoxy, haloalkyl, cyano, CHO, CO2H, alkoxycarbonyl, (substituted) Ph, PhO, PhS, PhCH2O, PhCH2S, etc.] were prepared Thus, 2-(4,6-dimethoxypyrimidin-2-yl)oxybenzoic acid, carbonyldiimidazole, and Me2NOH were stirred 12 h in THF/CH2C12 to give title compound II. Several I at 0.31 g/a preemergent gave complete control of velvetleaf.

IT 135564-62-4P

(preparation of, as herbicide)

RN 135564-62-4 HCAPLUS

CN Methanamine, N-[[2-[(4,6-dimethoxy-2-pyrimidinyl)oxy]benzoyl]oxy]-N-methyl- (9CI) (CA INDEX NAME)

EP 135280

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IC
    ICM C07D239-34
    ICS C07D239-52; C07D239-38; C07D401-12; C07D213-62; C07C239-22;
         C07C323-62; A01N043-54
CC
    28-16 (Heterocyclic Compounds (More Than One Hetero Atom))
    Section cross-reference(s): 5
ΙT
    135564-62-4P
                  135564-63-5P
                                 135564-64-6P
                                                 135564-65-7P
    135564-66-8P 135564-67-9P 135564-68-0P
    135564-69-1P 135564-70-4P 135564-71-5P
    135564-72-6P 135564-73-7P 135564-74-8P
    135564-75-9P 135564-76-0P 135564-77-1P
    135564-78-2P 135564-79-3P 135564-80-6P
    135564-81-7P
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    135564-84-0P
                  135564-85-1P
                                  135564-86-2P
                                               135564-87-3P
    135564-88-4P
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                                                 135564-91-9P
    135564-92-0P
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                                 135564-98-6P
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    135565-00-3P
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    135565-04-7P 135565-05-8P 135565-06-9P
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    135565-10-5P 135565-11-6P 135565-12-7P 135565-13-8P
    135587-60-9P
        (preparation of, as herbicide)
ΙT
    135565-14-9P 135565-15-0P 135565-16-1P
    135565-17-2P 135565-18-3P 135565-19-4P
    135565-20-7P
                  135565-21-8P
                                 135565-22-9P 135565-23-0P
    135565-24-1P
                  135565-25-2P
        (preparation of, as herbicide intermediate)
L68 ANSWER 46 OF 46 HCAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:
                        1985:221335 HCAPLUS Full-text
DOCUMENT NUMBER:
                        102:221335
ORIGINAL REFERENCE NO.: 102:34745a,34748a
TITLE:
                        Free radical polymerization and the produced
                        polymers
INVENTOR(S):
                        Solomon, David Henry; Rizzardo, Ezio; Cacioli,
PATENT ASSIGNEE(S):
                        Commonwealth Scientific and Industrial Research
                        Organization, Australia
SOURCE:
                        Eur. Pat. Appl., 63 pp.
                        CODEN: EPXXDW
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
    PATENT NO.
                        KIND
                               DATE
                                           APPLICATION NO.
                                                                  DATE
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19850327

Α2

EP 1984-304756

19840711

EP	135280			А3	19860	0813						
EP	135280			В1	1993	0512						
	R: B	BE, CH,	DE,	FR,	GB, IT,	LI,	NL					
AU	843037	8		Α	19850	0117		AU	1984-30378		198307	11
AU	571240			В2	1988	0414						
JP	600894	:52		Α	1985	0520		JΡ	1984-144097		198407	11
JP	050065	37		В	1993	0126						
US	458142	:9		Α	1986	0408		US	1984-629929		198407	11
PRIORIT	Y APPLN	I. INFO	).:					AU	1983-223	A	198307	11
								AU	1983-224	A	198307	11
								AU	1983-225	A	198307	11
								AU	1984-3578	A	198402	10

OTHER SOURCE(S): MARPAT 102:221335

ED Entered STN: 29 Jun 1985

AB Fifteen compds. R1R2R3CN(OX)CR4R5R6 (X = a group having ≥1 C and is such that the free radical formed by cleavage of the O-X bond is capable of polymerizing unsatd. monomers; R1, R2, R5, and R6 = alkyl or substituted alkyl of sufficient length to provide steric hindrance; R3 and R4 = alkyl, substituted alkyl, or a group which completes a cyclic structure containing the N atom) are prepared and used as initiators for the polymerization, including block and graft polymerization, of unsatd. monomers. The initiators permit control of the growth steps of polymns. to produce oligomeric polymers. Thus, a solution of 400 mg di-tert-butyl nitroxide and 234 mg di-tert-Bu peroxyoxalate [1876-22-8] in 5 mL styrene [100-42-5] was heated 2 h at 50° to prepare N-(2-tert-butyl-1- phenylethoxy)-N,N-di-tert-butylamine (I) [96724-60-6]. A solution of 52 mg I in 3 mL Me acrylate was heated 0.5 h at 100° to prepare 0.42 g polymer Bu3OCH2CHPh[CH2CH(CO2Me)]250N(CMe3)2.

IT 96724-56-0P

(preparation of, as radical polymerization catalyst)

RN 96724-56-0 HCAPLUS

CN Piperidine, 1-[2-(1,1-dimethylethoxy)-1-methyl-1-phenylethoxy]-2,6-dimethyl-2,6-dipropyl- (CA INDEX NAME)

$$\begin{array}{c|c}
 & Ph \\
 & C - CH_2 - OBu - t \\
 & Me \\
 & Me \\
 & Pr - n
\end{array}$$

IC ICM C08F004-00

ICS C08F291-00

CC 35-3 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 27

IT 96724-56-0P 96724-57-1P 96724-58-2P

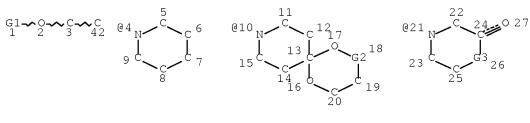
96724-59-3P 96724-60-6P

(preparation of, as radical polymerization catalyst)

=> d que 169

1 SEA FILE=HCAPLUS ABB=ON PLU=ON US20050215691/PN

L5 STR





VAR G1=4/10/21/34/37

REP G2=(0-1) CH2

VAR G3=O/N

NODE ATTRIBUTES:

NSPEC IS RC AT 42 NSPEC IS RC AT 43 DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 43

STEREO ATTRIBUTES: NONE

L9 STR

VAR G1=4/10/21/34/37

VAR G3=O/N VAR G4=46/40 NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

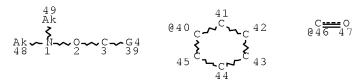
NUMBER OF NODES IS 47

STEREO ATTRIBUTES: NONE

L11 7221 SEA FILE=REGISTRY SSS FUL L9

L14 694 SEA FILE=REGISTRY SUB=L11 SSS FUL L5

L17 STR



VAR G4=46/40 NODE ATTRIBUTES: CONNECT IS E1 RC AT 48 CONNECT IS E1 RC AT 49 DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 14

STEREO ATTRIBUTES: NONE

L19 240 SEA FILE=REGISTRY SUB=L11 SSS FUL L17

L21 STR

VAR G1=4/10/21/34

REP G2=(0-1) CH2

VAR G3=O/N

VAR G4=46/40

## NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

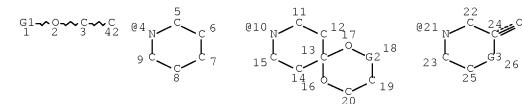
#### GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 44

#### STEREO ATTRIBUTES: NONE

L23	2091	SEA FILE=REGISTRY SUR	B=L11 SSS FU	L L21	
L24	262	SEA FILE=HCAPLUS ABB=	ON PLU=ON	L14	
L25	144	SEA FILE=HCAPLUS ABB=	ON PLU=ON	L19	
L29	209	SEA FILE=HCAPLUS ABB=	ON PLU=ON	L24(L)PREP/RL	
L32	74	SEA FILE=HCAPLUS ABB=	ON PLU=ON	L29 AND (PLASTIC?	OR
		POLYMER?)/SC,SX			
L33	60	SEA FILE=HCAPLUS ABB=	ON PLU=ON	L32 AND (INITIAT?	OR
		CATALYST? OR ACTIVAT?	?)		
L34	58	SEA FILE=HCAPLUS ABB=	ON PLU=ON	L33 NOT L19	
T.35		STR			





VAR G1=4/10/21/34

REP G2=(0-1) CH2

VAR G3=O/N

NODE ATTRIBUTES:

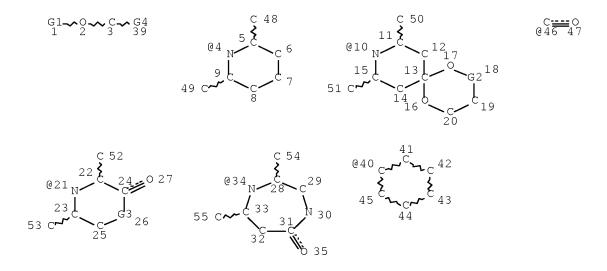
NSPEC IS RC AT 42 NSPEC IS RC AT 43 DEFAULT MLEVEL IS ATOM DEFAULT ECLEVEL IS LIMITED

#### GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED NUMBER OF NODES IS 40

#### STEREO ATTRIBUTES: NONE

L37	198	SEA	FILE=REGISTR	Y SUB=L1	4 SSS FU	L35	ō	
L38	96	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L37		
L39	40	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L34	AND	L38
L42		STR						



VAR G1=4/10/21/34

REP G2=(0-1) CH2

VAR G3=O/N

VAR G4=46/40

NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

#### GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 52

#### STEREO ATTRIBUTES: NONE

L44	1677	SEA FILE=REGISTRY	SUB=L23 SSS FUI	L L42
L45	614	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L44
L46	399	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L45(L)PREP/RL
L47	271	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L46 AND (PLASTIC? OR
		POLYMER?)/SC,SX		
L48	208	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L47 AND (INITIAT? OR
		CATALYST? OR ACTIV.	AT?)	
L49	1	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L48 AND L1
L50	2	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L48 AND CLAY?
L51	4	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L48 AND POF/RL
L52	174	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L48 AND CAT/RL
L53	6	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L52 AND MOA/RL
L54	15	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L52 AND (PEP OR PYP)/RL
L55	22	SEA FILE=HCAPLUS A	BB=ON PLU=ON	(L49 OR L50 OR L51) OR
		(L53 OR L54)		
L56	13	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L25 AND L46
L57	35	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L55 OR L56
L58	6	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L39 AND ALKOXYAMINE
		INITIATOR?		
L59	40	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L39 OR L58
L60	5	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L57 AND ALKOXYAMINE
		INITIATOR?		
L61	35	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L57 OR L60
L62	26	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L45 AND ALKOXYAMINE
		INITIATOR?		
L63	26	SEA FILE=HCAPLUS A	BB=ON PLU=ON	L62 AND (PLASTIC? OR

D(	٦T	.VI	MER	21	/SC.	SX

L64	56	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L61 OR L63
L65	6	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L38 AND ALKOXYAMINE
		INI	TIATOR?			
L66	40	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L59 OR L65
L67	10	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L64 AND L66
L69	30	SEA	FILE=HCAPLUS	ABB=ON	PLU=ON	L66 NOT L67

=> sel 169 hit rn 1-

E257 THROUGH E342 ASSIGNED

=> d 169 1-30 ibib ed abs hitstr hitind

L69 ANSWER 1 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2007:379556 HCAPLUS Full-text

DOCUMENT NUMBER: 146:411508

TITLE: Positive-working resist composition containing

cationic sulfonium salt and method of forming

pattern

INVENTOR(S): Hirano, Shuji; Kawanishi, Yasuhiro PATENT ASSIGNEE(S): Fuji Photo Film Co., Ltd., Japan SOURCE: Jpn. Kokai Tokkyo Koho, 57pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2007086516	A	20070405	JP 2005-276396	20050922
PRIORITY APPLN. INFO.:			JP 2005-276396	20050922

OTHER SOURCE(S): MARPAT 146:411508

ED Entered STN: 05 Apr 2007

GΙ

AB Disclosed is a pos.-working resist composition comprising (a) a polymer with  $\geq 3$  polymer chains via  $\geq 1$  branch whose solubility increases upon interaction with an acid and (b) a cationic sulfonium salt represented by I (R1-13 = H, substituent; and Z = single or divalent bonding group). The use of the cationic sulfonium salt improved line edge roughness when an electron beam was employed.

IT 933054-46-7P

(preparation of polymer for pos.-working resist composition)

RN 933054-46-7 HCAPLUS

CN Piperidine, 1,1',1'',1'''-[1,2,4,5-benzenetetrayltetrakis[methyleneoxy [1-[(1-ethoxyethoxy)phenyl]-2,1-ethanediyl]oxy]]tetrakis[2,2,6,6-tetramethyl- (CA INDEX NAME)

PAGE 1-A

O— CH— Me OEt PAGE 2-A

Me-CH-O
OEt

CC 74-5 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)

Section cross-reference(s): 35, 38, 76

IT Polymerization catalysts

(preparation of polymer for pos.-working resist composition)

IT 933054-46-7P

(preparation of polymer for pos.-working resist composition)

L69 ANSWER 2 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2006:1063135 HCAPLUS Full-text

DOCUMENT NUMBER: 145:419613

TITLE: Process for the preparation of star polymers via

epoxy-functionalized alkoxyamines

INVENTOR(S): Fink, Jochen; Pfaendner, Rudolf; Kramer, Andreas;

Fuso, Francesco

PATENT ASSIGNEE(S): Ciba Specialty Chemicals Holding Inc., Switz.

SOURCE: PCT Int. Appl., 23pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

	PATENT NO.					KIND DATE			APPLICATION NO.						DATE			
	WO	2006	1060	48		A1 20061012			1012	WO 2006-EP61045						20060327		
		W:	ΑE,	AG,	AL,	ΑM,	ΑT,	ΑU,	ΑZ,	BA,	BB,	BG,	BR,	BW,	BY,	BZ,	CA,	
			CH,	CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	
			GB,	GD,	GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KE,	KG,	KM,	
			KN,	KP,	KR,	KΖ,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	LY,	MA,	MD,	MG,	
			MK,	MN,	MW,	MX,	MZ,	NA,	NG,	NI,	NO,	NZ,	OM,	PG,	PH,	PL,	PT,	
			RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SM,	SY,	ΤJ,	TM,	TN,	TR,	TT,	
			TZ,	UA,	UG,	US,	UZ,	VC,	VN,	YU,	ZA,	ZM,	ZW					
		RW:	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	
			ΙE,	IS,	IT,	LT,	LU,	LV,	MC,	NL,	PL,	PT,	RO,	SE,	SI,	SK,	TR,	
			BF,	ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	TD,	
			TG,	BW,	GH,	GM,	KE,	LS,	MW,	MZ,	NA,	SD,	SL,	SZ,	TZ,	UG,	ZM,	
								KZ,										
	CA	2601	693			A1		2006	1012		CA 2	006-	2601	693		2	0060327	
	EP	1866	343			A1		2007	1219		EP 2	006-	7253	15		2	0060327	
		R:	ΑT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	
			ΙE,	IS,	IT,	LI,	LT,	LU,	LV,	MC,	NL,	PL,	PT,	RO,	SE,	SI,	SK, TR	
PRIOF	RIT	Y APP	LN.	INFO	.:					·	EP 2	005-	1026	21		A 2	0050404	

ED Entered STN: 12 Oct 2006

GΙ

AB A process for the preparation of star polymers, which are prepared by controlled free radical polymerization using epoxy-functionalized alkoxyamines and multifunctional compds. capable of reacting with the epoxy group. Thus, reaction of 0.01269 mol epoxy alkoxyamine I 9 h at 90° with 0.00423 mol tris(aminomethyl)amine in PhMe and polymerization of 90 g styrene 6 h at 130° in the presence of 2.5 g resulting multifunctional initiator gave star polymer.

WO 2006-EP61045

W 20060327

IT 912266-01-4P

(initiator; preparation of star polymers via initiators based on adducts of epoxy-functionalized alkoxyamines)

RN 912266-01-4 HCAPLUS

CN 2-Propanol, 1,1',1''-[nitrilotris(2,1-ethanediylimino)]tris[3-[4-[1-

 $\begin{tabular}{ll} [(3,3,8,8,10,10-hexamethyl-1,5-dioxa-9-azaspiro[5.5]undec-9-yl)oxy]ethyl]phenoxy]- (9CI) & (CA INDEX NAME) \end{tabular}$ 

PAGE 1-A

PAGE 1-B

— ОН

PAGE 2-B

$$-\mathsf{CH}_2-\mathsf{CH}_2-\mathsf{NH}-\mathsf{CH}_2-\mathsf{CH}-\mathsf{CH}_2-\mathsf{O} - \mathsf{Me} -$$

CC 35-3 (Chemistry of Synthetic High Polymers)

ST star shaped vinyl polymer epoxy alkoxyamine adduct initiator; trisaminomethylamine glycidyloxyphenylethoxyhexa methyldioxyazaspiroundecyloxy adduct initiator styrene polymn; controlled free radical polymn initiator vinyl compd

IT Adhesives

(preparation of star polymers via initiators based on adducts of epoxy-functionalized alkoxyamines for adhesives)

IT Crosslinking agents

(preparation of star polymers via initiators based on adducts of epoxy-functionalized alkoxyamines for crosslinkers)

IT Emulsifying agents

(preparation of star polymers via initiators based on adducts of epoxy-functionalized alkoxyamines for emulsifiers)

IT Ionomers

(preparation of star polymers via initiators based on adducts of epoxy-functionalized alkoxyamines for ionomers)

IT Polymer blend compatibilizers

(preparation of star polymers via initiators based on adducts of epoxy-functionalized alkoxyamines for polymer blend compatibilizers)

IT Surfactants

(preparation of star polymers via initiators based on adducts of epoxy-functionalized alkoxyamines for surfactants)

IT Polymerization

(radical, controlled; preparation of star polymers via initiators based on adducts of epoxy-functionalized alkoxyamines)

IT 4097-89-6, Tris(2-aminoethyl)amine 434898-80-3 (initiator precursor; preparation of star polymers via initiators based on adducts of epoxy-functionalized alkoxyamines)

IT 912266-01-4P

(initiator; preparation of star polymers via initiators based on adducts of epoxy-functionalized alkoxyamines)

IT 9003-53-6P, Polystyrene

(star; preparation of star polymers via initiators based on adducts of epoxy-functionalized alkoxyamines)

REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L69 ANSWER 3 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2006:146521 HCAPLUS Full-text

DOCUMENT NUMBER: 144:351070

TITLE: Photoresponsive poly(methyl methacrylate)2-

(polystyrene) 2 miktoarm star copolymer containing

an azobenzene moiety at the core

AUTHOR(S): Erdogan, Tuba; Gungor, Eda; Durmaz, Hakan; Hizal,

Gurkan; Tunca, Umit

CORPORATE SOURCE: Department of Chemistry, Istanbul Technical

University, Maslak, 34469, Turk.

SOURCE: Journal of Polymer Science, Part A: Polymer

Chemistry (2006), 44(4), 1396-1403

CODEN: JPACEC; ISSN: 0887-624X

PUBLISHER: John Wiley & Sons, Inc.

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 16 Feb 2006

AB We prepared a novel miktoarm star copolymer with an azobenzene unit at the core via combination of atom transfer radical polymerization (ATRP) and nitroxide-mediated free radical polymerization (NMP) routes. For this purpose, first, mikto-functional initiator, 3, with tertiary bromide (for ATRP) and 2,2,6,6-tetramethylpiperidin-1-yloxy (TEMPO) (for NMP) functionalities and an azobenzene moiety at the core was synthesized. The initiator 3 thus obtained was used in the subsequent living radical polymerization routes such as ATRP of MMA and NMP of St, resp., to give A2B2 type miktoarm star copolymer, (PMMA)2-(PSt)2 with an azobenzene unit at the core with controlled mol. weight and low polydispersity (Mw/Mn < 1.15). The photoresponsive properties of 3 and (PMMA)2-(PSt)2 miktoarm star copolymer were investigated.

IT 881388-21-2P

(initiator; photoresponsive poly(Me methacrylate)2-

(polystyrene)2 miktoarm star copolymer containing an azobenzene moiety at core)

RN 881388-21-2 HCAPLUS

CN Benzoic acid, 4,4'-azobis-, bis[2-[(2-bromo-2-methyl-1-oxopropoxy)methyl]-2-methyl-3-oxo-3-[2-phenyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethoxy]propyl] ester (9CI) (CA INDEX NAME)

PAGE 1-B

IT 881388-22-3P

(macroinitiator; photoresponsive poly(Me methacrylate)2(polystyrene)2 miktoarm star copolymer containing an azobenzene moiety
at core)

RN 881388-22-3 HCAPLUS

CN Benzoic acid, 4,4'-azobis-, bis[2-[(2-bromo-2-methyl-1-oxopropoxy)methyl]-2-methyl-3-oxo-3-[2-phenyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethoxy]propyl] ester, telomer with methyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 881388-21-2

CMF C66 H86 Br2 N4 O14

PAGE 1-B

CM 2

CRN 9011-14-7

CMF (C5 H8 O2)x

CCI PMS

CM 3

CRN 80-62-6

CMF C5 H8 O2

IT 881388-23-4P

(star polymer; photoresponsive poly(Me methacrylate)2-(polystyrene)2 miktoarm star copolymer containing an azobenzene moiety at core)

RN 881388-23-4 HCAPLUS

CN Benzoic acid, 4,4'-azobis-, bis[2-[(2-bromo-2-methyl-1-oxopropoxy)methyl]-2-methyl-3-oxo-3-[2-phenyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethoxy]propyl] ester, telomer with ethenylbenzene and methyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 881388-21-2

CMF C66 H86 Br2 N4 O14

PAGE 1-B

CM 2

CRN 25034-86-0

CMF (C8 H8 . C5 H8 O2) $\times$ 

CCI PMS

CM 3

CRN 100-42-5 CMF C8 H8

 $H_2C \longrightarrow CH - Ph$ 

CM 4

CRN 80-62-6 CMF C5 H8 O2



CC 35-4 (Chemistry of Synthetic High Polymers)

IT Polymerization

Polymerization catalysts

(atom transfer, living, radical; photoresponsive poly(Me methacrylate)2-(polystyrene)2 miktoarm star copolymer containing an azobenzene moiety at core)

IT 881388-21-2P

(initiator; photoresponsive poly(Me methacrylate)2-

(polystyrene)2 miktoarm star copolymer containing an azobenzene moiety at core)

IT 881388-22-3P

(macroinitiator; photoresponsive poly(Me methacrylate)2(polystyrene)2 miktoarm star copolymer containing an azobenzene moiety
at core)

IT 3030-47-5, Pentamethyldiethylenetriamine

(polymerization catalyst ligand; photoresponsive poly(Me methacrylate)2-(polystyrene)2 miktoarm star copolymer containing an azobenzene moiety at core)

IT 7758-89-6, Copper (I) chloride

(polymerization catalyst; photoresponsive poly(Me methacrylate)2-(polystyrene)2 miktoarm star copolymer containing an azobenzene moiety at core)

IT 10252-29-6, 4,4'-Bis(chlorocarbonyl)azobenzene 778603-97-7 (reactant in initiator preparation; photoresponsive poly(Me methacrylate)2-(polystyrene)2 miktoarm star copolymer containing an azobenzene moiety at core)

IT 881388-23-4P

(star polymer; photoresponsive poly(Me methacrylate)2-(polystyrene)2 miktoarm star copolymer containing an azobenzene moiety at core)

REFERENCE COUNT: 29 THERE ARE 29 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L69 ANSWER 4 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2006:41766 HCAPLUS <u>Full-text</u>

DOCUMENT NUMBER: 144:293151

TITLE: Syntheses of AB2 3- and AB4 5-miktoarm star copolymers by combination of the anionic

ring-opening polymerization of

hexamethylcyclotrisiloxane and nitroxide-mediated

radical polymerization of styrene

AUTHOR(S): Miura, Yozo; Yamaoka, Keisuke; Mannan, Md. Abdul CORPORATE SOURCE: Department of Applied Chemistry, Graduate School

of Engineering, Osaka City University, Osaka,

558-8585, Japan

SOURCE: Polymer (2006), 47(2), 510-519

CODEN: POLMAG; ISSN: 0032-3861

PUBLISHER: Elsevier Ltd.

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 17 Jan 2006

AB 
AB2 3- and AB4 5-microarm star copolymers were prepared by combination of the anionic ring-opening polymerization (AROP) of hexamethylcyclotrisiloxane (D3) and the TEMPO-mediated radical polymerization of styrene (St). Initially, two kinds of dendritic multifunctional initiators were prepared One has a 4-bromobutoxy group and two TEMPO-based alkoxyamines and the other has a 4-bromobutoxy group and four TEMPO-based alkoxyamines. Treatment of the multifunctional initiators with tert-butyllithium gave the corresponding lithiobutoxy derivs., and AROP of D3 by the lithiobutoxy derivs. gave poly(D3) with M w/Mn of 1.07-1.12. Nitroxide-mediated radical polymerization of St by the poly(D3)s at 120 °C gave AB2 3- and AB4 5-arm star copolymers with Mw/Mn of 1.15-1.28. Their structures were analyzed by means of 1H NMR and SEC measurements.

IT 878806-51-0P 878806-53-2P

(syntheses of star copolymers by combination of anionic ring-opening polymerization of hexamethylcyclotrisiloxane and nitroxide-mediated radical polymerization of styrene)

RN 878806-51-0 HCAPLUS

CN Lithium, [4-[3,5-bis[[4-[1-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl]phenyl]methoxy]phenoxy]butyl]- (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 878806-53-2 HCAPLUS

CN Lithium, [4-[3,5-bis[[3,5-bis[[4-[1-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl]phenyl]methoxy]phenyl]methoxy]phenoxy]butyl]-(CA INDEX NAME)

PAGE 1-A

Me Me Me 
$$CH_2$$
  $CH_2$   $CH_2$ 

PAGE 1-B

PAGE 2-A

Me---

PAGE 2-B

$$\frac{1}{\sqrt{N}} \int_{\mathbb{R}^{N}} Me$$

IT 878806-48-5P 878806-49-6P 878806-50-9P 878806-52-1P

(syntheses of star copolymers by combination of anionic ring-opening polymerization of hexamethylcyclotrisiloxane and nitroxide-mediated radical polymerization of styrene)

RN 878806-48-5 HCAPLUS

CN Phenol, 3,5-bis[[4-[1-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl]phenyl]methoxy]- (CA INDEX NAME)

PAGE 1-A

$$\begin{array}{c} \text{Me} & \text{Me} \\ \text{N} & \text{O-CH} \\ \text{Me} \end{array}$$

PAGE 1-B

CN

RN 878806-49-6 HCAPLUS

Phenol, 3,5-bis[[3,5-bis[[4-[1-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl]phenyl]methoxy]phenyl]methoxy]- (CA INDEX NAME)

## PAGE 1-B

RN

CN Piperidine, 1,1'-[[5-(4-bromobutoxy)-1,3-phenylene]bis(oxymethylene-4,1-phenyleneethylideneoxy)]bis[2,2,6,6-tetramethyl- (9CI) (CA INDEX NAME)

PAGE 1-B

RN 878806-52-1 HCAPLUS

CN Piperidine, 1,1',1'',1'''-[[5-(4-bromobutoxy)-1,3-phenylene]bis[oxymethylene-5,1,3-benzenetriylbis(oxymethylene-4,1-phenyleneethylideneoxy)]]tetrakis[2,2,6,6-tetramethyl-(9CI) (CA INDEX NAME)

Me Me 
$$CH_2$$
  $CH_2$   $C$ 

PAGE 1-B

- CC 35-4 (Chemistry of Synthetic High Polymers)
- ST hexamethylcyclotrisiloxane styrene star copolymer prepn ring opening polymn; nitroxide mediated radical polymn styrene star copolymer; dendrimer initiator prepn ring opening
- IT Polymerization

Polymerization catalysts

(anionic, ring-opening; syntheses of star copolymers by combination of anionic ring-opening polymerization of hexamethylcyclotrisiloxane and nitroxide-mediated radical polymerization of styrene)

IT Polymerization

Polymerization catalysts

(radical; syntheses of star copolymers by combination of anionic ring-opening polymerization of hexamethylcyclotrisiloxane and nitroxide-mediated radical polymerization of styrene)

IT 878806-51-0P 878806-53-2P

(syntheses of star copolymers by combination of anionic ring-opening polymerization of hexamethylcyclotrisiloxane and

nitroxide-mediated radical polymerization of styrene)

IT 878806-48-5P 878806-49-6P 878806-50-9P

878806-52-1P

(syntheses of star copolymers by combination of anionic ring-opening polymerization of hexamethylcyclotrisiloxane and nitroxide-mediated radical polymerization of styrene)

REFERENCE COUNT:

THERE ARE 46 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L69 ANSWER 5 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2005:574020 HCAPLUS Full-text

46

DOCUMENT NUMBER: 144:254458

TITLE: Synthesis of A3B3-type polystyrene-poly(methyl

methacrylate) miktoarm star polymers via combination of stable free radical and atom

transfer radical polymerization routes

AUTHOR(S): Durmaz, Hakan; Aras, Sermin; Hizal, Gurkan; Tunca,

Umit

CORPORATE SOURCE: Department of Chemistry, Istanbul Technical

University, Istanbul, 34469, Turk.

SOURCE: Designed Monomers and Polymers (2005), 8(3),

203-210

CODEN: DMPOF3; ISSN: 1385-772X

PUBLISHER: VSP
DOCUMENT TYPE: Journal
LANGUAGE: English
ED Entered STN: 04 Jul 2005

AB An A3B3-type miktoarm star polymer was prepared utilizing a "core-out" method via combination of stable free radical polymerization (SFRP) and atom transfer radical polymerization (ATRP). First, SFRP of styrene was carried out by using a miktofunctional initiator, benzene-1,3,5-tricarboxylic acid tris{3-(2-bromo-2-methyl- propionyloxy)-2-methyl-2-[2-phenyl-2-(2,2,6,6-tetramethylpiperidin-1- yloxy)ethoxy-carbonyl]propyl} ester, at 125°C. Second, previously obtained polystyrene (A3-type PSt) precursor with three bromine functionalities in the core was used as a macroinitiator for ATRP of Me methacrylate (MMA) in the presence of Cu(I)Cl and pentamethyldiethylenetriamine at 90°C in order to give an A3B3-type miktoarm

pentamethyldiethylenetriamine at  $90^{\circ}\text{C}$  in order to give an A3B3-type miktoars star polymer, (PSt)3(-PMMA)3, with controlled mol. weight and moderate polydispersity (Mw/Mn < 1.41).

IT 876931-41-8P

(core and multifunctional initiator; synthesis of A3B3-type polystyrene-poly(Me methacrylate) miktoarm star polymers via combination of stable free radical and atom transfer radical polymerization routes)

RN 876931-41-8 HCAPLUS

CN 1,3,5-Benzenetricarboxylic acid, tris[2-[(2-bromo-2-methyl-1-oxopropoxy)methyl]-2-methyl-3-oxo-3-[2-phenyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethoxy]propyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

PAGE 2-A

- CC 35-4 (Chemistry of Synthetic High Polymers)
- IT Polymerization catalysts

(radical; synthesis of A3B3-type polystyrene-poly(Me methacrylate) miktoarm star polymers via combination of stable free radical and atom transfer radical polymerization routes)

IT 876931-41-8P

(core and multifunctional initiator; synthesis of

A3B3-type polystyrene-poly(Me methacrylate) miktoarm star polymers via combination of stable free radical and atom transfer radical polymerization routes)

ΤТ 9003-53-6P, Polystyrene

> (three-arms star, multifunctional initiator; synthesis of A3B3-type polystyrene-poly(Me methacrylate) miktoarm star polymers via combination of stable free radical and atom transfer radical polymerization routes)

REFERENCE COUNT:

THERE ARE 14 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L69 ANSWER 6 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2005:217463 HCAPLUS Full-text

14

DOCUMENT NUMBER: 142:411785

TITLE: Synthesis of AB2 3- and AB4 5-miktoarm star

copolymers initiated from dendritic triand penta-functional initiators by

combination of ring-opening polymerization of

 $\epsilon$ -caprolactone and nitroxide-mediated radical polymerization of styrene

AUTHOR(S): Miura, Yozo; Sakai, Yumi; Yamaoka, Keisuke

CORPORATE SOURCE: Department of Applied Chemistry, Graduate School

of Engineering, Osaka City University, Osaka,

558-8585, Japan

SOURCE: Macromolecular Chemistry and Physics (2005),

206(4), 504-512

CODEN: MCHPES; ISSN: 1022-1352 Wiley-VCH Verlag GmbH & Co. KGaA

DOCUMENT TYPE: Journal LANGUAGE: English Entered STN: 11 Mar 2005

AΒ Well-defined AB2 3- and AB4 5-miktoarm star copolymers were prepared by combination of ring-opening polymerization (ROP) and nitroxide-mediated radical polymerization (NMRP) using dendritic tri- and penta-functional initiators. Initially, two kinds of dendritic initiators having one benzylic OH and two or four TEMPO-based alkoxyamine moieties were prepared Using them, ROP of  $\varepsilon$ -caprolactone was carried out at room temperature to give poly-( $\varepsilon$ caprolactone)s carrying two or four alkoxyamine moieties. NMRP of styrene from the poly(ε-caprolactone)s was carried out at 120°C to give AB2 3- and AB4 5-miktoarm star copolymers, which were analyzed by 1H NMR and SEC. The Mns increased linearly with conversion and the Mw/Mns were in the range 1.10-1.37, showing that well-defined AB2 3- and AB4 5-miktoarm star copolymers were formed.

492446-77-2 ΙT

PUBLISHER:

(preparation of dendritic penta-functional initiators for synthesis of AB4 5-miktoarm star copolymers)

492446-77-2 HCAPLUS RN

CN Benzenemethanol, 3,5-bis[[4-[1-[(2,2,6,6-tetramethyl-1piperidinyl)oxy]ethyl]phenyl]methoxy]- (CA INDEX NAME)

PAGE 1-A

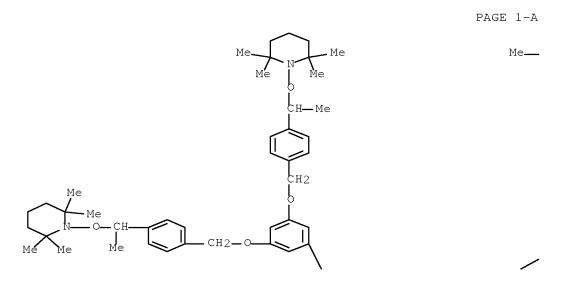
PAGE 1-B

IT 850538-80-6P

(preparation of dendritic penta-functional initiators for synthesis of AB4 5-miktoarm star copolymers)

RN 850538-80-6 HCAPLUS

CN Benzenemethanol, 3,5-bis[[3,5-bis[[4-[1-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl]phenyl]methoxy]phenyl]methoxy]- (CA INDEX NAME)



PAGE 1-B

IT 492446-78-3

(preparation of dendritic penta-functional initiators for synthesis of AB4 5-miktoarm star copolymers)

RN 492446-78-3 HCAPLUS

CN Piperidine, 1,1'-[[5-(bromomethyl)-1,3-phenylene]bis(oxymethylene-4,1-phenyleneethylideneoxy)]bis[2,2,6,6-tetramethyl- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B



CC 35-7 (Chemistry of Synthetic High Polymers)

IT Polyesters, preparation

(lactone-based; synthesis of AB2 3- and AB4 5-miktoarm star copolymers initiated from dendritic tri- and

penta-functional initiators)

IT Polymerization catalysts

(radical; synthesis of AB2 3- and AB4 5-miktoarm star copolymers initiated from dendritic tri- and penta-functional initiators)

IT Polymers, preparation

(star-branched; synthesis of AB2 3- and AB4 5-miktoarm star copolymers initiated from dendritic tri- and penta-functional initiators)

IT 209550-24-3 492446-77-2

(preparation of dendritic penta-functional initiators for synthesis of AB4 5-miktoarm star copolymers)

IT 850538-80-6P

(preparation of dendritic penta-functional initiators for synthesis of AB4 5-miktoarm star copolymers)

IT 29654-55-5, 3,5-Dihydroxybenzyl alcohol 492446-78-3 (preparation of dendritic penta-functional initiators for synthesis of AB4 5-miktoarm star copolymers)

24980-41-4P, ε-Caprolactone homopolymer 25248-42-4P,

 $\varepsilon$ -Caprolactone homopolymer, sru 29010-44-4P,

 $\epsilon$ -Caprolactone-styrene copolymer

(synthesis of AB2 3- and AB4 5-miktoarm star copolymers initiated from dendritic tri- and penta-functional initiators)

REFERENCE COUNT:

CORPORATE SOURCE:

ΙT

32 THERE ARE 32 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L69 ANSWER 7 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2005:15040 HCAPLUS <u>Full-text</u>

DOCUMENT NUMBER: 142:261830

TITLE: Multi-armed, TEMPO-functionalized unimolecular

initiators for starburst dendrimer

synthesis via stable free radical polymerization.

2. Tris (1,3,5)benzyloxy unimers

AUTHOR(S): Ghani, Mohmad Asri Abd; Abdallah, Dalia; Kazmaier,

Peter M.; Keoshkerian, Barkev; Buncel, Erwin Department of Chemistry, Queen's University,

Kingston, ON, K7L 3N6, Can.

SOURCE: Canadian Journal of Chemistry (2004), 82(9),

1403-1412

CODEN: CJCHAG; ISSN: 0008-4042

PUBLISHER: National Research Council of Canada

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 142:261830

ED Entered STN: 07 Jan 2005

The synthesis of the trifunctionalized TEMPO-modified unimol. initiators, unimers I, II, and III is described. Unimer I was prepared via an SN2 type Williamson ether coupling of 1,3,5-tris(iodomethyl)benzene with a TEMPO-containing ethylbenzene hydroxy derivative. The synthesis of unimer II, however, was accomplished through SN1 reaction of 1,3,5-tris(bromomethyl)benzene with the hydroxy-ethylbenzene TEMPO derivative in the presence of silver triflate. Synthesis of unimer III started from phloroglucinol and an SNAr reaction with 1-fluoro-4-nitrobenzene, followed by reduction to the amino compound and Schiff base formation with the TEMPO-derivatized aromatic aldehyde. Stable free radical polymerization (SFRP) of styrene and acetoxystyrene with unimer I are also described with mol. wts. and polydispersities reported. It is concluded that the SFRP of styrene with a triradical initiator meets the requirements of a living system.

IT 372522-45-7P 845745-22-4P 845745-23-5P

(preparation of multi-armed, TEMPO-functionalized trisbenzyloxyunimol. initiators for radical polymerization of styrene and acetoxystyrene)

RN 372522-45-7 HCAPLUS

CN Piperidine, 1,1',1''-[1,3,5-benzenetriyltris[methyleneoxy(1-phenyl-2,1-ethanediyl)oxy]]tris[2,2,6,6-tetramethyl- (9CI) (CA INDEX NAME)

RN 845745-22-4 HCAPLUS

CN Piperidine, 1,1',1''-[1,3,5-benzenetriyltris(methyleneoxymethylene-4,1-phenyleneethylideneoxy)]tris[2,2,6,6-tetramethyl- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

PAGE 2-A

RN 845745-23-5 HCAPLUS

CN Benzenamine, 4,4',4''-[1,3,5-benzenetriyltris(oxy)]tris[N-[[4-[1-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl]phenyl]methylene]- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

PAGE 2-A

- CC 35-3 (Chemistry of Synthetic High Polymers)
- ST styrene acetoxystyrene radical polymn catalyst multiarmed TEMPO functionalized trisbenzyloxyunimol
- IT Molecular weight

Molecular weight distribution

(preparation of multi-armed, TEMPO-functionalized trisbenzyloxyunimol. initiators for radical polymerization of styrene and acetoxystyrene)

IT Polymerization catalysts

(radical; preparation of multi-armed, TEMPO-functionalized trisbenzyloxyunimol. initiators for radical polymerization of styrene and acetoxystyrene)

IT Polymers, preparation

(star-branched; preparation of multi-armed, TEMPO-functionalized trisbenzyloxyunimol. initiators for radical polymerization of styrene and acetoxystyrene)

IT 9003-53-6P, Polystyrene 24979-78-0DP, Poly(p-acetoxystyrene), hydrolyzed 24979-78-0P, Poly(p-acetoxystyrene) 188203-34-1P, p-Acetoxystyrene-styrene block copolymer

(3-armed; preparation of multi-armed, TEMPO-functionalized trisbenzyloxyunimol. initiators for radical polymerization of styrene and acetoxystyrene)

IT 209550-24-3P 372522-45-7P 845745-22-4P 845745-23-5P

(preparation of multi-armed, TEMPO-functionalized trisbenzyloxyunimol. initiators for radical polymerization of styrene and acetoxystyrene)

IT 94-36-0, BPO, reactions 100-41-4, Ethylbenzene, reactions 100-42-5, Styrene, reactions 108-73-6, 1,3,5-Trihydroxybenzene 350-46-9, 4-Fluoro-nitrobenzene 1074-61-9, 4-Vinylbenzyl alcohol 1876-22-8, Di-tert-butylperoxyoxalate 2564-83-2, TEMPO 18226-42-1, 1,3,5-Tris(bromomethyl)benzene 90678-60-7, 1,3,5-Tris(iodomethyl)benzene 102852-91-5 209550-23-2 (preparation of multi-armed, TEMPO-functionalized trisbenzyloxyunimol. initiators for radical polymerization of styrene and acetoxystyrene)

IT 81913-53-3P 102852-92-6P 154554-67-3P 161776-41-6P (preparation of multi-armed, TEMPO-functionalized trisbenzyloxyunimol. initiators for radical polymerization of styrene and acetoxystyrene)

REFERENCE COUNT: 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L69 ANSWER 8 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2005:15039 HCAPLUS Full-text

DOCUMENT NUMBER: 142:261829

TITLE: Multi-armed, TEMPO-functionalized unimolecular

initiators for starburst dendrimer

synthesis via stable free radical polymerization.

1. Tri azofunctionalized unimer

AUTHOR(S): Abdallah, Dalia; Ghani, Mohmad Asri Abd;

Cunningham, Michael F.; Kazmaier, Peter M.;

Keoshkerian, Barkev; Buncel, Erwin

CORPORATE SOURCE: Department of Chemistry, Queen's University,

Kingston, ON, K7L 3N6, Can.

SOURCE: Canadian Journal of Chemistry (2004), 82(9),

1393-1402

CODEN: CJCHAG; ISSN: 0008-4042 National Research Council of Canada

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 142:261829

ED Entered STN: 07 Jan 2005

PUBLISHER:

The synthesis of azobenzene-functionalized multi-armed unimol. initiators or "unimers" that can be polymerized using styrene or styrenic derivs. via TEMPO (2,2,6,6-tetramethylpiperidenyl-1-oxyl) mediated stable free radical polymerization (SFRP) is described. The unimers are composed of an azobenzene-functionalized core and a TEMPO-modified unit. Homopolymers and

copolymers of styrene and acetoxystyrene were synthesized using the mono-and trifunctionalized unimers as initiators under bulk conditions with average mol. wts. and polydispersities reported. The studies lay the groundwork for further investigations involving SFRP towards building a light harvesting system by introducing chromophores onto the polymer chains for capturing light and thence transferring it to the azobenzene core.

IT 845728-32-7P 845728-34-9P

(multi-armed, TEMPO-functionalized unimol. initiators for starburst dendrimer synthesis via stable radical polymerization)

RN 845728-32-7 HCAPLUS

CN Piperidine, 1,1'-[1,2-ethanediylbis(4,1-phenyleneazo-4,1-phenyleneoxymethylene-4,1-phenyleneethylideneoxy)]bis[2,2,6,6-tetramethyl-(9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 845728-34-9 HCAPLUS

CN Piperidine, 1,1',1''-[1,3,5-benzenetriyltris(oxy-4,1-phenyleneazo-4,1-phenyleneoxymethylene-4,1-phenyleneethylideneoxy)]tris[2,2,6,6-tetramethyl- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 2-B

**—** Ме

CC

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ST
    triazofunctionalized unimer initiator radical polymn styrene
     acetoxystyrene dendrimer synthesis; TEMPO modified styrene
     acetoxystyrene copolymer polystyrene polyacetoxystyrene dendrimer
     synthesis
ΙT
    Molecular weight
     Molecular weight distribution
        (multi-armed, TEMPO-functionalized unimol. initiators for
        starburst dendrimer synthesis via stable radical polymerization)
     Polymerization catalysts
ΙT
        (radical; multi-armed, TEMPO-functionalized unimol.
        initiators for starburst dendrimer synthesis via stable
        radical polymerization)
ΙT
    Polymers, preparation
        (star-branched; multi-armed, TEMPO-functionalized unimol.
        initiators for starburst dendrimer synthesis via stable
        radical polymerization)
     24979-78-0P, Poly(p-acetoxystyrene)
ΤT
                                          188203-34-1P,
     p-Acetoxystyrene-styrene block copolymer
        (linear or 3-armed; multi-armed, TEMPO-functionalized unimol.
        initiators for starburst dendrimer synthesis via stable
        radical polymerization)
     1876-22-8P, Di-tert-Butyl peroxyoxalate
TΤ
                                              845728-31-6P
     845728-32-7P 845728-34-9P
        (multi-armed, TEMPO-functionalized unimol. initiators for
        starburst dendrimer synthesis via stable radical polymerization)
ΙT
     9003-53-6P, Polystyrene 24979-78-0DP, Poly(p-acetoxystyrene),
     hydrolyzed 53746-03-5P, p-Acetoxystyrene-styrene copolymer
        (multi-armed, TEMPO-functionalized unimol. initiators for
        starburst dendrimer synthesis via stable radical polymerization)
TΤ
     75-91-2, tert-Butylhydroperoxide 79-37-8, Oxalyl chloride
     108-73-6, 1,3,5-Benzenetriol 108-95-2, Phenol, reactions
     621-95-4
              1689-82-3 2564-83-2, TEMPO
                                             4748-78-1
                                                         7632-00-0,
     Sodium nitrite
        (multi-armed, TEMPO-functionalized unimol. initiators for
        starburst dendrimer synthesis via stable radical polymerization)
ΙT
     768-59-2P
               57825-30-6P 90264-99-6P
                                           102852-91-5P
                                                          102852-92-6P
     492446-76-1P
                  845728-30-5P
                                  845728-33-8P
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35-3 (Chemistry of Synthetic High Polymers)

(multi-armed, TEMPO-functionalized unimol. initiators for

starburst dendrimer synthesis via stable radical polymerization) IT 558-13-4

(multi-armed, TEMPO-functionalized unimol. initiators for

starburst dendrimer synthesis via stable radical polymerization)

REFERENCE COUNT: 33 THERE ARE 33 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L69 ANSWER 9 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2004:76619 HCAPLUS Full-text

DOCUMENT NUMBER: 140:112181

TITLE: Manufacture of polymers having dissociative

electron attachment groups and scission of the

polymer main chains

INVENTOR(S): Ichikawa, Tsuneki; Koizumi, Hitoshi; Shimizu,

Akira

PATENT ASSIGNEE(S): Shin-Etsu Chemical Industry Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 10 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2004026981	A	20040129	JP 2002-184301	20020625
PRIORITY APPLN. INFO.:			JP 2002-184301	20020625

ED Entered STN: 30 Jan 2004

The polymers, useful for resists for radiation lithog, are manufactured by reaction of monomers capable of living polymerization with compds. having living polymerization initiating groups on both ends via dissociative electron attachment groups which cleave by attachment of dissociative electron. The polymer main chain is cut by irradiation of electromagnetic wave or particle beam having energy sufficient to ionize the mol. Thus, 4-[1-(2,2,6,6- tetramethylpiperidinyl-1-oxy)ethyl]benzyl alc. was esterified with 4-[1-(2,2,6,6- tetramethylpiperidinyl-1-oxy)ethyl]benzoic acid, then the resulting ester was added 2% to styrene and heated to 130°, when living radical polymerization reaction occurred. The polymer was cut in half by irradn of  $\gamma$ -ray.

IT 647849-32-9P

(manufacture of living polymers having radiation-cleavable structure in main chain)

RN 647849-32-9 HCAPLUS

$$\begin{array}{c} \text{Me} & \text{Me} \\ \text{N} & \text{O-CH} \\ \end{array}$$

IC ICM C08F004-00 ICS C08F012-00; C08F020-00; C08J003-28; C08L101-00

CC 37-3 (Plastics Manufacture and Processing) Section cross-reference(s): 25, 27, 74

IT 647849-32-9P

(manufacture of living polymers having radiation-cleavable structure in main chain)

L69 ANSWER 10 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2004:54261 HCAPLUS <u>Full-text</u>

DOCUMENT NUMBER: 140:94477

TITLE: Initiators for nitroxide-mediated

polymerization based on azlactone or their

ring-opened derivatives

INVENTOR(S): Fansler, Duane D.; Lewandowski, Kevin M.;

Wendland, Michael S.; Gaddam, Babu N.

PATENT ASSIGNEE(S): 3M Innovative Properties Company, USA

SOURCE: U.S., 11 pp.
CODEN: USXXAM

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PAT	TENT	NO.			KINI	)	DATE			APPL	ICAT	ION :	NO.			ATE 
US	6680	 362			B1		2004	0120		 US 2	003-	 3587	67			0030205
US	2004	1528.	53		A1		2004	0805		US 2	003-	7269	56		2	0031203
US	6784	265			В2		2004	0831								
WO	2004	0721	39		A1		2004	0826		WO 2	004-1	US11	30		2	0040116
WO	2004	0721	39		A8		2005	0224								
	W:	ΑE,	AG,	AL,	ΑM,	AT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BW,	BY,	BZ,	CA,
		CH,	CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,
		GB,	GD,	GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	KΕ,	KG,	KP,
		KR,	KΖ,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,
		MX,	MZ,	NA,	NI											
	RW:	BW,	GH,	GM,	ΚE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	AT,
		BE,	BG,	CH,	CY,	CZ,	DE,	DK,	EE,	ES,	FI,	FR,	GB,	GR,	HU,	IE,
		ΙΤ,	LU,	MC,	NL,	PT,	RO,	SE,	SI,	SK,	TR,	BF,	ВJ,	CF,	CG,	CI,
		CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,	ΝE,	SN,	TD,	ΤG				
EP	1590	387			A1		2005	1102		EP 2	004-	7029	49		2	0040116
	R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	ΙΤ,	LI,	LU,	NL,	SE,	MC,
		PT,	IE,	SI,	LT,	LV,	FΙ,	RO,	MK,	CY,	AL,	TR,	BG,	CZ,	EE,	HU, SK
JP	2006	5166	69		Τ		2006	0706		JP 2	006-	5028	58		2	0040116
PRIORITY	Y APP	LN.	INFO	.:						US 2	003-	3587	67	ì	A3 2	0030205
										WO 2	004-	US11	30	Ţ	W 2	0040116

OTHER SOURCE(S): MARPAT 140:94477

ED Entered STN: 22 Jan 2004

AB A controlled radical polymerization initiator comprises R5ZCO(CH2)nCR3R4NHCOQCR1[(CH2)qX][CH2CR1[QCONHCR3R4(CH2)nCOZR5]]mONR22, wherein X is an H, an alkyl group, a cycloalkyl group, a heterocyclic group, an arenyl group, an aryl group, a nitrile, an acyl group or the residue of a free-radical initiator; R1 is H, an alkyl group, a cycloalkyl group, a heterocyclic group, an arenyl group or an aryl group; ON(R2)2 is the residue of an organonitroxide; R3 and R4 are each independently selected from an alkyl, a cycloalkyl group, an aryl group, an arenyl group, or R3 and R4 taken together with the carbon to which they are attached form a carbocyclic ring; Q is a linking group selected from a covalent bond, (CH2)o, CO2(CH2)o,

CO2(CH2CH2O)o, CONR6(CH2)o, COS(CH2)o, where o is 1 to 12, and R6 is H, an alkyl group, a cycloalkyl group, an arenyl group, a heterocyclic group or an aryl group; each n is 0 or 1; m is 0 to 20; q is 0 or 1; Z is O, S or NR6, wherein R6 is H, an alkyl group, a cycloalkyl group, an arenyl group, a heterocyclic group or an aryl group; R5 is an organic or inorg. moiety and has a valency of p. The initiators have an azlactone or ring-opened azlactone moiety to provide telechelic (co)polymers. 4,4-Dimethyl-2-[1-(2,2,6,6-tetramethylpiperidin-1-yloxy)-ethyl]-4H-ox azol-5-one was prepared and used in polymerization of styrene.

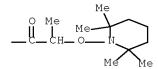
IT 642479-67-2P

(initiators for nitroxide-mediated polymerization based on azlactone or their ring-opened derivs.)

RN 642479-67-2 HCAPLUS

CN Propanamide, N,N',N''-(nitrilotri-2,1-ethanediyl)tris[2-methyl-2-[[1-oxo-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]propyl]amino]- (9CI) (CA INDEX NAME)

PAGE 2-B



ICM C08F002-00 IC INCL 526217000; 526222000; 526224000; 526265000; 526271000; 526287000; 526291000; 526303100; 526304000; 526306000 CC 35-3 (Chemistry of Synthetic High Polymers) azlactone initiator nitroxide mediated radical polymn ST TΤ Polymerization catalysts (controlled radical; initiators for nitroxide-mediated polymerization based on azlactone or their ring-opened derivs.) 642479-67-2P ΤТ (initiators for nitroxide-mediated polymerization based on azlactone or their ring-opened derivs.) 642479-65-0P ΤТ (initiators for nitroxide-mediated polymerization based on azlactone or their ring-opened derivs.) 9003-53-6P, Styrene homopolymer ΤТ (initiators for nitroxide-mediated polymerization based on azlactone or their ring-opened derivs.) 642479-61-6P ΙT 642479-59-2P 642479-63-8P (initiators for nitroxide-mediated polymerization based on azlactone or their ring-opened derivs.) TΤ 62-57-7, 2-Aminoisobutyric acid 563-76-8, 2-Bromopropionyl bromide 2564-83-2, TEMPO 4097-89-6, Tris(2-aminoethyl)amine 21055-93-6. Sodium diethylthiocarbamate (initiators for nitroxide-mediated polymerization based on azlactone or their ring-opened derivs.) REFERENCE COUNT: 21 THERE ARE 21 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L69 ANSWER 11 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2004:42401 HCAPLUS Full-text

DOCUMENT NUMBER: 140:236159

TITLE: Synthesis of Nanosized "Cored" Star Polymers

AUTHOR(S): Beil, James B.; Zimmerman, Steven C.

CORPORATE SOURCE: Department of Organic Chemistry, University of

Illinois at Urbana Champaign, Urbana, IL, 61801,

USA

SOURCE: Macromolecules (2004), 37(3), 778-787

CODEN: MAMOBX; ISSN: 0024-9297

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 19 Jan 2004

AB A synthetic approach to nanosized "cored" star polymers is reported. A 5,10,15,20-tetrakis(4-carboxyphenyl)porphyrin core was functionalized with four 2,2,6,6-tetramethylpiperidinyl-1-oxy (TEMPO) initiating groups. Four-armed star copolymers of styrene and 4-hydroxystyrene were synthesized and functionalized with 3,5-di(3-buten-1-oxy) benzyl bromide groups but exhibited poor solubility As an alternative, 5,10,15,20-tetrakis(3',5'-

dihydroxyphenyl)porphyrin was functionalized with 2-bromo-2-methyl-propionyl groups capable of initiating atom transfer radical polymerization (ATRP). Copolymn. of the core initiator with 1-but-3-enyl-4-vinylbenzene and styrene at low conversion produced soluble eight-armed star block copolymers. Through the ring-closing metathesis (RCM) reaction, the alkene groups of the polymer were intramolecularly cross-linked. The ester groups linking the cross-linked polymer arms to the porphyrin core were hydrolyzed, producing a "cored" star polymer with a mol. weight of approx. 20 kDa and a polydispersity index (PDI) of 1.5.

IT 668420-45-9P

(polymerization catalyst; in preparation of nanosized "Cored" star polymers)

RN 668420-45-9 HCAPLUS

CN Benzoic acid, 4,4',4'',4'''-(21H,23H-porphine-5,10,15,20-tetrayl)tetrakis-, tetrakis[2-phenyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl] ester (9CI) (CA INDEX NAME)

Me **\** 

PAGE 2-B

PAGE 3-A

CC 35-4 (Chemistry of Synthetic High Polymers)

ST Star block copolymer TEMPO contg core initiator ATRP

IT 668420-45-9P 668420-51-7P

(polymerization catalyst; in preparation of nanosized "Cored" star polymers)

IT 245107-65-7DP, crosslinked derivative by metathesis catalyst, and hydrolyzed

(preparation of nanosized "Cored" star polymers)

IT 245107-65-7P 668420-49-3DP, derivative with a core initiator, hydrolyzed, reaction product with di-substituted benzyl bromide 668420-50-6DP, derivative with styrene-hydroxy styrene copolymer

(star polymer; preparation of nanosized "Cored" star polymers)

REFERENCE COUNT: 50 THERE ARE 50 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L69 ANSWER 12 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2004:33987 HCAPLUS Full-text

DOCUMENT NUMBER: 140:94467

TITLE: Azlactone initiators for

nitroxide-mediated polymerization

INVENTOR(S): Lewandowski, Kevin M.; Fansler, Duane D.;

Wendland, Michael S.; Gaddam, Babu N. 3M Innovative Properties Company, USA

PATENT ASSIGNEE(S): 3M Innovative SOURCE: U.S., 11 pp.

CODEN: USXXAM

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT 1	NO.			KIN	D i	DATE		1	APPL	ICAT	ION I	NO.		D.	ATE
00 00 1110					B1 20040113 A1 20040805					003- 003-		20030205 20031203			
US 6784 WO 2004		27		B2 A1		2004 2004		1	WO 2	003-	US41.	366		2	0031223
₩:	CH,	CN,	CO,	CR,	CU,	AU, CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,
	KR,	KZ,	LC,	LK,	LR,	HR, LS, OM,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,
	YU,	ZA,	ZM,	ZW	·	TM,	·	ŕ	ŕ	·	ŕ	ŕ	ŕ	ŕ	·
RW:	AZ,	BY,	KG,	KZ,	MD,	MW, RU, GB,	ТJ,	TM,	AT,	BE,	BG,	CH,	CY,	CZ,	DE,
	SE,	SI,		TR,	BF,	BJ,									•

AU	2003	2975	40		A1		2004	0906		ΑU	200	3-2	2975.	40		2	20031	223
EP	1590	374			A1		2005	1102		ΕP	200	3-8	31592	23		2	20031	223
EP	1590	374			В1		2006	0614										
	R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GF	R, I	Γ,	LI,	LU,	NL,	SE,	MC,	
		PT,	IE,	SI,	LT,	LV,	FI,	RO,	MK,	C.7	7, A	L,	TR,	BG,	CZ,	EE,	HU,	SK
JP	2006	5141	33		Τ		2006	0427		JΡ	200	4 - 5	5683	43		2	20031	223
AT	3299	33			Τ		2006	0715		ΑT	200	3-8	31592	23		2	20031	223
PRIORIT	Y APP	LN.	INFO	.:						US	200	3 – 3	35872	24	i	A3 2	20030	205
										MO	200	3 <b>—</b> T	1941	366	Ţ	M 2	0031	223

OTHER SOURCE(S): MARPAT 140:94467

ED Entered STN: 15 Jan 2004

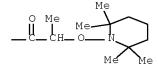
AB The initiators have an azlactone or ring-opened azlactone moiety to provide telechelic (co)polymers. AzTEMPO (0.00066 mol) and styrene (0.132 mol) were mixed in toluene, the solution was deoxygenated by bubbling N (g) through it for 30 min and heated to 130°, after 16 h the product had Mn 20,611 g/mol, then a 1% solution of tris(2-aminoethyl)amine (0.000226 mol) in toluene was added in two portions to give a three-arm polystyrene of Mn 50,061 g/mol. IT 642479-67-2P

(azlactone initiators for nitroxide-mediated polymerization of styrene)

RN 642479-67-2 HCAPLUS

CN Propanamide, N,N',N''-(nitrilotri-2,1-ethanediyl)tris[2-methyl-2-[[1-oxo-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]propyl]amino]- (9CI) (CA INDEX NAME)

PAGE 2-B



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ICM C08F002-00
TC
INCL 526204000; 526217000; 526222000; 526224000; 526265000; 526271000;
     526287000; 526291000; 526303100; 526304000
     35-3 (Chemistry of Synthetic High Polymers)
CC
     Section cross-reference(s): 67
     Polymerization catalysts
ΙT
        (radical; azlactone initiators for nitroxide-mediated
        polymerization of styrene)
ΙT
     642479-67-2P
        (azlactone initiators for nitroxide-mediated polymerization of
        styrene)
ΙT
     9003-53-6DP, Polystyrene, azlactone-terminated
        (azlactone initiators for nitroxide-mediated polymerization of
        styrene)
ΙT
     2564-83-2, TEMPO 29513-26-6, 2-Vinyl-4,4-dimethyl azlactone
        (azlactone initiators for nitroxide-mediated polymerization of
        styrene)
TΤ
     4097-89-6, Tris(2-aminoethyl)amine
        (coupling agent; azlactone initiators for
        nitroxide-mediated polymerization of styrene)
ΙT
     642479-65-0P
        (initiator; azlactone initiators for
        nitroxide-mediated polymerization of styrene)
ΙT
     642479-59-2P
        (intermediate; azlactone initiators for
        nitroxide-mediated polymerization of styrene)
ΙT
     642479-61-6P
        (reaction with TEMPO; azlactone initiators for
        nitroxide-mediated polymerization of styrene)
     563-76-8, 2-Bromopropionyl bromide
ΙT
        (reaction with aminoisobutyric acid; azlactone initiators
        for nitroxide-mediated polymerization of styrene)
ΙT
     62-57-7, 2-Aminoisobutyric acid
        (reaction with bromopropionyl bromide; azlactone initiators
        for nitroxide-mediated polymerization of styrene)
ΙT
     21055-93-6, Sodium diethylthiocarbamate
        (reaction with bromopropionylamino methylpropionic acid; azlactone
        initiators for nitroxide-mediated polymerization of styrene)
     642479-63-8P
TT
        (ring closure reaction; azlactone initiators for
        nitroxide-mediated polymerization of styrene)
     9003-53-6P, Polystyrene
ΙT
        (star, 3-arm; azlactone initiators for nitroxide-mediated
        polymerization of styrene)
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REFERENCE COUNT: 21 THERE ARE 21 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L69 ANSWER 13 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2003:42249 HCAPLUS Full-text

DOCUMENT NUMBER: 138:107153

TITLE: Multifunctional alkoxyamines based on

polyalkylpiperidines, polyalkylpiperazinones and

polyalkylmorpholinones and their use as polymerization regulators/initiators

INVENTOR(S): Kramer, Andreas; Muehlebach, Andreas; Nesvadba,

Peter; Zink, Marie-Odile; Hintermann, Tobias

PATENT ASSIGNEE(S): Ciba Specialty Chemicals Holding Inc., Switz.

SOURCE: PCT Int. Appl., 48 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PA	TENT	NO.			KIN	)	DATE			APPL	ICAT	ION :	NO.		D.	ATE
WO	2003	0044	 71		A1	_	2003	0116		 WO 2	002-	EP71	 31		2	0020627
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		CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	ES,	FI,	GB,	GD,
		GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	ΚE,	KG,	KP,	KR,	KΖ,
		LC,	LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,
		NO,	NZ,	OM,	PH,	PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	ΤJ,
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		CH,	CY,	DE,	DK,	ES,	FΙ,	FR,	GB,	GR,	ΙE,	ΙT,	LU,	MC,	NL,	PT,
		SE,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,	NE,
		SN,	TD,	ΤG												
CA	2450	232			A1		2003	0116		CA 2	002-	2450	232		2	0020627
AU	2002	3252	79		A1		2003	0121		AU 2	002-	3252	79		2	0020627
EP	1417	171			A1		2004	0512		EP 2	002-	7582	82		2	0020627
EP	1417	171			В1		2007	1121								
	R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	ΙT,	LI,	LU,	NL,	SE,	MC,
		PT,	ΙE,	SI,	LT,	LV,	FI,	RO,	MK,	CY,	AL,	TR				
CN	1522	247			Α		2004	0818		CN 2	002-	8134	56		2	0020627
	2005															0020627
MX	2003	PA11	833		Α		2004	0326		MX 2	003-	PA11	833		2	0031218
US	2004	1673	03		A1		2004	0826		US 2	003-	4825	46		2	0031230
US	6936	670			В2		2005	0830								
PRIORIT	Y APP	LN.	INFO	.:						EP 2	001-	8106	64		A 2	0010705
										WO 2	002-	EP71	31	,	W 2	0020627

OTHER SOURCE(S): MARPAT 138:107153

ED Entered STN: 17 Jan 2003

AB The instant invention relates to multifunctional alkoxyamines based on polyalkylpiperidines, polyalkylpiperazinones and polyalkylmorpholinones and their use as polymerization regulators/initiators. Further subjects of the invention are a polymerizable composition comprising an ethylenically unsatd. monomer (e.g., styrene) or oligomer and the alkoxyamine compound as well as a process for polymerization and a process for preparation of the compds.

IT 485844-67-5P 485844-69-7P 485844-70-0P 485844-71-1P 485844-72-2P 485844-74-4P 485844-75-5P 485844-77-7P 485844-78-8P

485844-79-9P 485844-80-2P 485844-81-3P

(multifunctional alkoxyamines based on polyalkylpiperidines, polyalkylpiperazinones and polyalkylmorpholinones and their use as polymerization regulators/initiators)

RN 485844-67-5 HCAPLUS

CN Propanoic acid, 2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]-, 1,2-ethanediyl ester (9CI) (CA INDEX NAME)

RN 485844-69-7 HCAPLUS

CN Propanoic acid, 2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]-, 1,4-cyclohexanediyl ester (9CI) (CA INDEX NAME)

RN 485844-70-0 HCAPLUS

CN Propanamide, N, N'-1, 6-hexanediylbis[2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]- (CA INDEX NAME)

RN 485844-71-1 HCAPLUS

CN Propanamide, N,N'-1,6-hexanediylbis[2-[(2,6-diethyl-2,3,6-trimethyl-4-oxo-1-piperidinyl)oxy]- (CA INDEX NAME)

RN 485844-72-2 HCAPLUS

CN Poly(oxy-1,2-ethanediyl),  $\alpha$ -[2-[(2,6-diethyl-2,3,6-trimethyl-4-oxo-1-piperidinyl)oxy]-1-oxopropyl]- $\omega$ -[2-[(2,6-diethyl-2,3,6-trimethyl-4-oxo-1-piperidinyl)oxy]-1-oxopropoxy]- (9CI) (CA INDEX NAME)

RN 485844-74-4 HCAPLUS

CN Poly(oxy-1,4-butanediyl),  $\alpha$ -[2-[(2,6-diethyl-2,3,6-trimethyl-4-oxo-1-piperidinyl)oxy]-1-oxopropyl]- $\omega$ -[2-[(2,6-diethyl-2,3,6-trimethyl-4-oxo-1-piperidinyl)oxy]-1-oxopropoxy]- (9CI) (CA INDEX NAME)

RN 485844-75-5 HCAPLUS

CN Propanoic acid, 2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]-, 2-[[2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]-1-oxopropoxy]methyl]-2-ethyl-1,3-propanediyl ester (9CI) (CA INDEX NAME)

RN 485844-77-7 HCAPLUS

CN Propanoic acid, 2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]-, 1,3,5-benzenetriyl ester (9CI) (CA INDEX NAME)

RN 485844-78-8 HCAPLUS

CN Propanoic acid, 2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]-, 2,2-bis[[2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]-1-oxopropoxy]methyl]-1,3-propanediyl ester (9CI) (CA INDEX NAME)

RN 485844-79-9 HCAPLUS

CN D-Glucitol, hexakis[2-[(2,6-diethyl-4-hydroxy-2,3,6-trimethyl-1-piperidinyl)oxy]propanoate] (9CI) (CA INDEX NAME)

Absolute stereochemistry.

RN 485844-80-2 HCAPLUS

CN Propanoic acid, 2-[[4-(1,1-dimethylethyl)-2,2-diethyl-6,6-dimethyl-3-oxo-1-piperazinyl]oxy]-, 1,2-ethanediyl ester (9CI) (CA INDEX NAME)

RN 485844-81-3 HCAPLUS

CN Propanoic acid, 2-[(3,3-diethyl-5,5-dimethyl-2-oxo-4-morpholinyl)oxy]-, 1,2-ethanediyl ester (9CI) (CA INDEX NAME)

IC ICM C07D211-94

ICS C08F004-00; C08F002-38; C08F012-08; C08F020-18

CC 35-3 (Chemistry of Synthetic High Polymers)

ST piperidine alkyl initiator polymn; piperazinone alkyl initiator polymn; morpholinone alkyl initiator polymn

IT Polymerization catalysts

(radical; multifunctional alkoxyamines based on
polyalkylpiperidines, polyalkylpiperazinones and
polyalkylmorpholinones and their use as polymerization regulators/
initiators)

IT 485844-67-5P 485844-69-7P 485844-70-0P

485844-71-1P 485844-72-2P 485844-74-4P

485844-75-5P 485844-77-7P 485844-78-8P

485844-79-9P 485844-80-2P 485844-81-3P

(multifunctional alkoxyamines based on polyalkylpiperidines, polyalkylpiperazinones and polyalkylmorpholinones and their use as polymerization regulators/initiators)

IT 3030-47-5 5468-93-9 51210-48-1 61745-37-7 200192-34-3

248603-09-0 264279-68-7 264279-93-8 286470-47-1 286470-49-3

300707-30-6 485844-68-6 485844-73-3 485844-76-6

(multifunctional alkoxyamines based on polyalkylpiperidines, polyalkylpiperazinones and polyalkylmorpholinones and their use as polymerization regulators/initiators)

REFERENCE COUNT:

THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L69 ANSWER 14 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN

5

ACCESSION NUMBER: 2002:964328 HCAPLUS Full-text

DOCUMENT NUMBER: 138:39710

TITLE: N-alkoxy 4-imino piperidine polymerization

regulators and their use in free radical-mediated polymerization of vinyl monomers to low dispersity

polymers

INVENTOR(S): Nesvadba, Peter; Hintermann, Tobias; Kramer,

Andreas; Zink, Marie-Odile; Bugnon, Lucienne

PATENT ASSIGNEE(S): Ciba Specialty Chemicals Holding Inc., Switz.

SOURCE: PCT Int. Appl., 51 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PA:	PATENT NO.					KIND		DATE		APPLICATION NO.						DATE	
WO	2002	10083	31		A1		2002	1219		WO 2	2002-	EP61	08		2	0020604	
	W:	ΑE,	AG,	AL,	AM,	AT,	ΑU,	AZ,	BA,	BB,	BG,	BR,	BY,	BZ,	CA,	CH,	
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		GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	JP,	ΚE,	KG,	KP,	KR,	KZ,	
		LC,	LK,	LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	
		NO,	NZ,	OM,	PH,	PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	ΤJ,	
		TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US,	UΖ,	VN,	YU,	ZA,	ZM,	ZW		
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		CH,	CY,	DE,	DK,	ES,	FΙ,	FR,	GB,	GR,	ΙE,	ΙT,	LU,	MC,	NL,	PT,	
		SE,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,	MR,	NE,	
		SN,	TD,	ΤG													
CA	2443	718			A1		2002	1219		CA 2	2002-	2443	718		2	0020604	
AU	2002	32880	06		A1		2002	1223		AU 2	2002-	3288	06		2	0020604	
EP	1397	349			A1		2004	0317		EP 2	2002-	7645	77		2	0020604	
	R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	ΙΤ,	LI,	LU,	NL,	SE,	MC,	
		PT,	ΙE,	SI,	LT,	LV,	FΙ,	RO,	MK,	CY,	AL,	TR					
	1514						2004				–				_	0020604	
	2005						2005	0127		JP 2	2003-	5036	00		2	0020604	
MX	20031	PA101	132		Α		2004	0310		MX 2	2003-	PA10	132		2	0031106	
US	2004	1765	53		A1		2004	0909		US 2	2003-	4801	88		2	0031209	
US	7199	245			В2		2007	0403									
RIORIT	Y APP:	LN.	INFO	.:						EP 2	2001-	8105	67	1	A 2	0010613	
										EP 2	2001-	8111	54		A 2	0011128	
										WO 2	2002-	EP61	08	1	W 2	0020604	

OTHER SOURCE(S): MARPAT 138:39710

ED Entered STN: 20 Dec 2002

AB The present invention relates to selected 4-imino-N-alkoxy-polyalkyl-piperidine compds. preparation, a polymerizable composition comprising a) at least one ethylenically unsatd. monomer and b) a 4-imino-N-alkoxy-polyalkyl-piperidine compound Further aspects of the present invention are a process for polymerizing ethylenically unsatd. monomers, and the use of 4-imino-N-alkoxy-polyalkyl-piperidine compds. for controlled polymerization The intermediate N-oxyl derivs., a composition of the N-oxyl derivs. with ethylenically unsatd. monomers and a free radical initiator, as well as a process for polymerization are also subjects of the present invention.

II 478697-58-4P

 $\hbox{ (preparation of $N-$alkoxy $4-$imino piperidine polymerization regulators and their}\\$ 

use in free radical-mediated vinyl monomer polymerization to low

dispersity polymers) RN 478697-58-4 HCAPLUS

CN 4-Piperidinone, 2,6-diethyl-2,3,6-trimethyl-1-(1-phenylethoxy)-, 0,0'-(1,6-dioxo-1,6-hexanediyl)dioxime (9CI) (CA INDEX NAME)

IC ICM C07D211-94

ICS C08F004-00

CC 35-3 (Chemistry of Synthetic High Polymers)

IT 478697-27-7P 478697-34-6P 478697-38-0P 478697-40-4P 478697-42-6P 478697-44-8P 478697-47-1P 478697-49-3P 478697-53-9P 478697-55-1P 478697-58-4P 478697-60-8P 478697-62-0P 478697-65-3P 478697-67-5P

(preparation of N-alkoxy 4-imino piperidine polymerization regulators and their

use in free radical-mediated vinyl monomer polymerization to low

dispersity polymers)
REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L69 ANSWER 15 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2002:868581 HCAPLUS Full-text

DOCUMENT NUMBER: 138:137635

TITLE: Synthesis of six-arm star polymer by

nitroxide-mediated "living" radical polymerization

AUTHOR(S): Miura, Yozo; Yoshida, Yuji

CORPORATE SOURCE: Department of Applied Chemistry, Graduate School

of Engineering, Osaka City University, Osaka,

558-8585, Japan

SOURCE: Polymer Journal (Tokyo, Japan) (2002), 34(10),

748-754

CODEN: POLJB8; ISSN: 0032-3896 Society of Polymer Science, Japan

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 15 Nov 2002

PUBLISHER:

AB A dendritic multifunctional initiator with six TEMPO-based alkoxyamine moieties was prepared from 4-bromoethylbenzene in seven steps. Six-arm star polymers were synthesized by the radical bulk polymerization of styrene using the dendritic alkoxyamine as an initiator. The styrene polymns. were carried out at 120° using the dendritic alkoxyamine concns. of 5.0, 12.8 and 18.8 mmol/L. When the alkoxyamine concentration was 5.0 mmol/L, the polydispersity of the resulting star polymers increased with conversion, and the polydispersity of the star polymer at 72% conversion was 1.59. When the alkoxyamine concns. were 12.8 and 18.8 mmol/L, the polymerization was well controlled to give star polymers with low polydispersity even at high conversions. Mol. weight of the star polymers determined by NMR significantly differed from GPC and light scattering detns. and was attributed to the unique structure of the polymers. The six-arm polystyrene synthesis showed features of the living polymerization with some side reactions at high monomer

conversion. The deviations from the living polymerization character were discussed on the basis of GPC elution curves of the star polymers.

IT 492446-77-2P 492446-78-3P

(in synthesis of multifunctional TEMPO-based radical initiators for production of six-arm star polymers)

RN 492446-77-2 HCAPLUS

CN Benzenemethanol, 3,5-bis[[4-[1-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl]phenyl]methoxy]- (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 492446-78-3 HCAPLUS

CN Piperidine, 1,1'-[[5-(bromomethyl)-1,3-phenylene]bis(oxymethylene-4,1-phenyleneethylideneoxy)]bis[2,2,6,6-tetramethyl- (9CI) (CA INDEX NAME)

PAGE 1-B

IT 492446-79-4P

(synthesis of multifunctional TEMPO-based radical initiators for production of six-arm star polymers)

- RN 492446-79-4 HCAPLUS
- CN Piperidine, 1,1',1'',1''',1'''',1''''-[1,3,5-benzenetriyltris[oxymethylene-5,1,3-benzenetriylbis(oxymethylene-4,1-phenyleneethylideneoxy)]]hexakis[2,2,6,6-tetramethyl- (9CI) (CA INDEX NAME)

PAGE 2-B

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CC
     35-3 (Chemistry of Synthetic High Polymers)
     Section cross-reference(s): 67
ST
     multifunctional radical living catalyst star polymer
     synthesis
     Polymerization
ΙT
        (living, radical, bulk; multifunctional TEMPO-based radical
        initiators for synthesis of six-arm star polymers)
ΙT
     Polymerization catalysts
        (living, radical; multifunctional TEMPO-based radical
        initiators for synthesis of six-arm star polymers)
     NMR (nuclear magnetic resonance)
ΙT
     Size-exclusion chromatography
        (mol. structure determination of six-arm star polymers produced using
        multifunctional TEMPO-based radical initiators)
     Coupling reaction
ΤТ
        (star-star coupling; synthesis of six-arm star polymers using
       multifunctional TEMPO-based radical initiators)
     Molecular structure-property relationship
ΙT
        (synthesis of six-arm star polymers using multifunctional
        TEMPO-based radical initiators)
     584-08-7, Potassium carbonate
ΤT
        (in synthesis of multifunctional TEMPO-based radical
        initiators for production of six-arm star polymers)
     492446-76-1P 492446-77-2P 492446-78-3P
ΤТ
        (in synthesis of multifunctional TEMPO-based radical
        initiators for production of six-arm star polymers)
     68-12-2, DMF, reactions 108-73-6, 1,3,5-Benzenetriol
                                                              558-13-4,
ΤТ
     Carbon tetrabromide 1585-07-5, 4-Bromoethylbenzene 2564-83-2,
            7789-60-8, Phosphorous tribromide 16940-66-2, Sodium
     borohydride
                 29654-55-5, 3,5-Dihydroxybenzyl alcohol
        (in synthesis of multifunctional TEMPO-based radical
        initiators for production of six-arm star polymers)
ΙT
     178625-97-3P
                   209550-23-2P
                                  209550-24-3P
        (in synthesis of multifunctional TEMPO-based radical
        initiators for production of six-arm star polymers)
     594-19-4, tert-Butyllithium 603-35-0, Triphenylphosphine, reactions
TT
     1876-22-8, Di-tert-butyl diperoxyoxalate
                                               17455-13-9, 18-Crown-6
        (in synthesis of multifunctional TEMPO-based radical
        initiators for production of six-arm star polymers)
     9003-53-6P, Polystyrene
ΙT
        (star, six-arm; multifunctional TEMPO-based radical
        initiators for synthesis of six-arm star polymers)
IΤ
     492446-79-4P
        (synthesis of multifunctional TEMPO-based radical
        initiators for production of six-arm star polymers)
REFERENCE COUNT:
                         27
                               THERE ARE 27 CITED REFERENCES AVAILABLE FOR
                               THIS RECORD. ALL CITATIONS AVAILABLE IN THE
                               RE FORMAT
L69 ANSWER 16 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:
                         2002:626615 HCAPLUS Full-text
DOCUMENT NUMBER:
                         137:338264
TITLE:
                         Synthesis of end-functionalized polymer with
                         cyclodextrin based on tempo-mediated radical
                         polymerization
AUTHOR(S):
                         Narumi, Atsushi; Miura, Yutaka; Satoh, Toshifumi;
                         Kaga, Harumi; Kakuchi, Toyoji
                         Div. Molecular Chem., Grad. Sch. Eng., Hokkaido
CORPORATE SOURCE:
                         Univ., Sapporo, 060-8628, Japan
```

SOURCE: Polymer Preprints (American Chemical Society,

Division of Polymer Chemistry) (2002), 43(2),

279-280

CODEN: ACPPAY; ISSN: 0032-3934

PUBLISHER: American Chemical Society, Division of Polymer

Chemistry

DOCUMENT TYPE: Journal; (computer optical disk)

LANGUAGE: English ED Entered STN: 20 Aug 2002

AB Polystyrene was end-functionalized with a cyclic oligosacchamide, cyclodextrin

(CD). First, mono-6-[4-[1'-(2'',2'',6'',6''-tetramethyl- 1''-

piperidinyloxy)ethyl]benzamido]-per-2,3,6-acetyl- $\beta$ - cyclodextrin was prepared and was used as initiator to polymerize styrene. The resulting polymer was deacetylated. A reversed-type micelle using the polymer with the  $\beta$ -CD core

was prepared

IT 474088-71-6P

(catalyst; in preparation of end-functionalized polymer with cyclodextrin based on tempo-mediated radical polymerization)

RN 474088-71-6 HCAPLUS

CN  $\beta$ -Cyclodextrin, 6A,6B,6C,6D,6E,6F-hexadeoxy-6A,6B,6C,6D,6E,6F-hexakis[[4-[1-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl]benzoyl]am ino]-, 2A,2B,2C,2D,2E,2F,2G,3A,3B,3C,3D,3E,3F,3G,6G-pentadecaacetate (9CI) (CA INDEX NAME)

PAGE 2-A

# PAGE 3-B

PAGE 5-A

CC = 35-4 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 44

IT Polymerization catalysts

(living, radical; in preparation of end-functionalized polymer with cyclodextrin based on tempo-mediated radical polymerization)

IT 474088-71-6P

(catalyst; in preparation of end-functionalized polymer with cyclodextrin based on tempo-mediated radical polymerization)

REFERENCE COUNT: 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE

RE FORMAT

L69 ANSWER 17 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2002:354020 HCAPLUS Full-text

DOCUMENT NUMBER: 136:370147

TITLE: Soluble polymer supports for organic synthesis

INVENTOR(S): Janda, Kim D.; Gravert, Dennis J. PATENT ASSIGNEE(S): The Scripps Research Institute, USA

SOURCE: U.S. Pat. Appl. Publ., 34 pp., Cont. of U.S. Ser.

No. 161,604. CODEN: USXXCO

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2002055124	A1	20020509	US 2001-996402	20011119
PRIORITY APPLN. INFO.:			US 1998-161604 A	1 19980923

ED Entered STN: 12 May 2002

GΙ

- AB Polymer supports for liquid-phase organic synthesis (LPOS) are employed in a process for transferring a chemical intermediate between immiscible solvents. These compds. are produced with an expanded range of solubility range in a variety of solvent systems. A sequence of normal and "living" free radical polymns. are employed to generate a library of block copolymers possessing either block or graft architecture with initiators having N:N and TEMPO groups tethered by ester or ether linkages for styrene, 4-tert-butylstyrene (I), 3,4-dimethoxystyrene, vinylpyrrolidinone, N-isopropylacrylamide, and 1-methacryloyloxy-2-phenyl-2-(2,2,6,6-tetramethyl-1-piperidinyloxy)ethane. A typical block copolymer was manufactured by polymerization of 7.75 mmol mg styrene 8 h at 70° in 1,2-dichlorobenzene in the presence of initiator II, and polymerization of 1.08 mmol I 12 h at 130° in the presence of 1.02 mg intermediate.
- IT 188119-33-7P 203382-60-9P 213994-38-8P

(polymerization initiator; soluble graft and block styrene (derivative)-based polymer supports for organic synthesis)

RN 188119-33-7 HCAPLUS

CN Pentanoic acid, 4,4'-azobis[4-cyano-, bis[2-phenyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl] ester (9CI) (CA INDEX NAME)

RN 203382-60-9 HCAPLUS

CN Pentanenitrile, 2,2'-azobis[2-methyl-5-[2-phenyl-2-[(2,2,6,6-

tetramethyl-1-piperidinyl)oxy]ethoxy]- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 213994-38-8 HCAPLUS

CN Pentanoic acid, 4,4'-azobis[4-cyano-, bis[2-[[4-[[(1,1-dimethylethoxy)carbonyl]amino]-2,2,6,6-tetramethyl-1-piperidinyl]oxy]-2-phenylethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

IC ICM G01N033-53

ICS G01N033-543; C08F008-30; C08F008-44; C08F008-32

INCL 435007100

CC 35-4 (Chemistry of Synthetic High Polymers)
 Section cross-reference(s): 23

ST soluble block graft polymer support combinatorial library org synthesis; methacryloyloxyphenyl tetramethylpiperidinyloxyethane

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polymer manuf; isopropylacrylamide block polymer manuf TEMPO azo
     initiator; vinylpyrrolidinone block polymer manuf TEMPO azo
     initiator; dimethoxystyrene block polymer manuf TEMPO azo
     initiator; styrene tertiary butylstyrene block copolymer manuf
     TEMPO azo initiator
ΙT
    Hydrogenation catalysts
        (polymer-supported; soluble graft and block styrene (derivative)-based
        polymer supports for organic synthesis)
ΙT
     108-55-4, Glutaric anhydride
                                    61478-29-3
        (hydrogenation catalyst precursor; soluble graft and block
        styrene (derivative)-based polymer supports for organic synthesis)
     12092-47-6
ΙT
        (hydrogenation datalyst; soluble graft and block styrene
        (derivative) - based polymer supports for organic synthesis)
     213994-93-5P
ΤT
        (hydrogenation catalyst; soluble graft and block styrene
        (derivative) - based polymer supports for organic synthesis)
                                   423126-07-2P
ΤТ
     213994-43-5P
                   213994-50-4P
        (polymerization initiator precursor; soluble graft and block
        styrene (derivative)-based polymer supports for organic synthesis)
     94-36-0, Benzoyl peroxide, reactions 100-42-5, Styrene, reactions
ΙT
                        2638-94-0, 4,4'-Azobis(cyanovaleric acid)
     2564-83-2, TEMPO
     14691-88-4, 4-Amino-TEMPO
                                17170-81-9, 4,4'-Azobis(4-cyanovaleroyl
     chloride)
                 24424-99-5, Di-tert-butyl dicarbonate
        (polymerization initiator precursor; soluble graft and block
        styrene (derivative)-based polymer supports for organic synthesis)
     188119-33-7P 203382-60-9P 213994-38-8P
ΙT
        (polymerization initiator; soluble graft and block styrene
        (derivative) - based polymer supports for organic synthesis)
ΙT
     161776-41-6
        (reactive polymerization initiator precursor; soluble graft and
        block styrene (derivative)-based polymer supports for organic synthesis)
ΙT
     213994-57-1P
        (reactive polymerization initiator; soluble graft and block styrene
        (derivative) - based polymer supports for organic synthesis)
L69 ANSWER 18 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:
                         2002:311344 HCAPLUS Full-text
                         137:79284
DOCUMENT NUMBER:
TITLE:
                         Syntheses of functional alkoxyamines and
                         application to syntheses of well-defined star
                         polymers
AUTHOR(S):
                         Miura, Yozo; Yoshida, Yuji
CORPORATE SOURCE:
                         Department of Applied Chemistry, Graduate School
                         of Engineering, Osaka City University, Osaka,
                         558-8585, Japan
SOURCE:
                         Macromolecular Chemistry and Physics (2002),
                         203(5/6), 879-888
                         CODEN: MCHPES; ISSN: 1022-1352
PUBLISHER:
                         Wiley-VCH Verlag GmbH
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
     Entered STN: 25 Apr 2002
ED
AB
     Three kinds of 2,2,6,6-tetramethylpiperidine-N-oxyl (TEMPO)-based
     alkoxyamines, 1-(4-iodophenyl)-(3), 1-(4-ethynylphenyl)-(4), and 1-[4-(1, 3, 4)]
     2-dio xaborinan-2-y1) phenyl]-1-(2,2,6,6-tetramethyl-1- piperidinyloxyl) ethanes
     (5) were prepared The Pd-catalyzed cross-coupling reaction of 3 with 1,3,5-
     triethynylbenzene or 1,3,5-tribromobenzene with 4 gave the corresponding
     1,3,5-tris(alkoxyaminophenylethynyl)benzene 11, and the Pd-catalyzed cross-
     coupling reaction of 5 with 1,3,5-tribromobenzenegave the corresponding 1,3,5-
```

tris(alkoxyaminophenyl)benzene 12. Bulk polymerization of styrene (St) at  $120\,^{\circ}\text{C}$  initiated with 11 and 12 were investigated. The first-order plots, linear relationships between and conversion, and low Mw/Mns of the formed poly(St) showed that the polymerization proceeded in the "living" fashion leading to formation of well-defined three-arm star polymers with Mw/Mn of 1.20-1.40.

IT 439904-91-3P 439904-92-4P 439904-93-5P 439904-94-6P

(polymerization initiator; syntheses of TEMPO-based functional alkoxyamines and their application to syntheses of well-defined star polystyrenes)

RN 439904-91-3 HCAPLUS

CN Piperidine, 1,1'-[1,3-phenylenebis(2,1-ethynediyl-4,1-phenyleneethylideneoxy)]bis[2,2,6,6-tetramethyl-(9CI) (CA INDEX NAME)

PAGE 1-A

$$\begin{array}{c} \text{Me} \\ \text{N} \\ \text{Me} \end{array} \begin{array}{c} \text{Me} \\ \text{CH} \\ \text{O} \end{array} \begin{array}{c} \text{C} \\ \text{CH} \\ \text{O} \end{array} \begin{array}{c} \text{Me} \\ \text{Me} \\ \text{O} \end{array}$$

PAGE 1-B

RN 439904-92-4 HCAPLUS

CN Piperidine, 1,1',1''-[1,3,5-benzenetriyltris(2,1-ethynediyl-4,1-phenyleneethylideneoxy)]tris[2,2,6,6-tetramethyl- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 439904-93-5 HCAPLUS

CN Piperidine, 1,1'-[1,4-phenylenebis(2,1-ethynediyl-4,1-phenyleneethylideneoxy)]bis[2,2,6,6-tetramethyl-(9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B



RN 439904-94-6 HCAPLUS

CN Piperidine, 1,1'-[[5'-[4-[1-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl]phenyl][1,1':3',1''-terphenyl]-4,4''-diyl]bis(ethylideneoxy)]bis[2,2,6,6-tetramethyl-(9CI) (CA INDEX NAME)

CC 35-3 (Chemistry of Synthetic High Polymers)

ST TEMPO based alkoxyamine synthesis styrene living polymn initiation

IT Amines, preparation

(alkoxylated, polymerization initiators; syntheses of TEMPO-based functional alkoxyamines and their application to syntheses of well-defined star polystyrenes)

IT Polymerization catalysts

(atom transfer, living, radical; syntheses of TEMPO-based functional alkoxyamines and their application to syntheses of well-defined star polystyrenes)

IT 154554-67-3

(polymerization initiator; syntheses of TEMPO-based functional alkoxyamines and their application to syntheses of well-defined star polystyrenes)

IT 439904-91-3P 439904-92-4P 439904-93-5P 439904-94-6P

(polymerization initiator; syntheses of TEMPO-based functional alkoxyamines and their application to syntheses of well-defined star polystyrenes)

REFERENCE COUNT:

THERE ARE 43 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L69 ANSWER 19 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2002:232041 HCAPLUS Full-text DOCUMENT NUMBER: 137:20656

43

JOCOMENI NOMBER. 137.20030

TITLE: Synthesis and characterization of star-shaped

polystyrene having  $\beta$ -cyclodextrin-core

AUTHOR(S): Sugimoto, Naoya; Matsuda, Takeshi; Miura, Yutaka;

Narumi, Atsushi; Kaga, Harumi; Kakuchi, Toyoji

CORPORATE SOURCE: Div. Molecular Chem., Grad. Sch. Eng., Hokkaido

Univ., Sapporo, 060-8628, Japan

SOURCE: Polymer Preprints (American Chemical Society,

Division of Polymer Chemistry) (2002), 43(1),

664-665

CODEN: ACPPAY; ISSN: 0032-3934

PUBLISHER: American Chemical Society, Division of Polymer

Chemistry

DOCUMENT TYPE: Journal; (computer optical disk)

LANGUAGE: English ED Entered STN: 27 Mar 2002

AB A cyclodextrin-based multifunctional initiator was prepared from amino-

cyclodextrin and N-(1-methyl-4-carbonylbenzyloxy)-2,2,6,6-

tetramethylpiperidine. Polymerization of styrene (St) was conducted by atom transfer radical polymerization using cyclodextrin-based multifunctional

initiator.

IT 434318-35-1P (initiator; in preparation and characterization of star-shaped

polystyrene having  $\beta$ -cyclodextrin-core)

RN 434318-35-1 HCAPLUS

CN  $\beta$ -Cyclodextrin, 6A,6B,6C,6D,6E,6F,6G-heptadeoxy-

6A, 6B, 6C, 6D, 6E, 6F, 6G-heptakis[[4-[1-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl]benzoyl]amino]- (9CI) (CA INDEX NAME)

PAGE 2-B

PAGE 3-B

CC = 35-4 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 44

ST cyclodextrin based multifunctional ATRP initiator prepn; star shaped polystyrene cyclodextrin core prepn

IT Polymerization

Polymerization catalysts

(atom transfer, radical; in preparation and characterization of star-shaped polystyrene having  $\beta\text{-cyclodextrin-core})$ 

IT 434318-35-1P

(initiator; in preparation and characterization of star-shaped polystyrene having  $\beta$ -cyclodextrin-core)

IT 9003-53-6DP, Poly(styrene), initiated by  $\beta\text{-cyclodextrin}$  based multifunctional TEMPO initiator

(star-shaped; synthesis and characterization of star-shaped polystyrene having  $\beta$ -cyclodextrin-core)

REFERENCE COUNT: 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE

#### RE FORMAT

L69 ANSWER 20 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2002:182192 HCAPLUS Full-text

DOCUMENT NUMBER: 136:217209

TITLE: Dual purpose electroactive copolymers, preparation

> thereof, and use in opto-electronic devices Hawker, Craig Jon; Klaerner, Gerrit; Lee,

Jeong-Ik; Lee, Victor Yee-Way; Miller, Robert

Dennis; Scott, John Campbell

PATENT ASSIGNEE(S): International Business Machines Corporation, USA;

The Board of Trustees of the Leland Stanford

Junior University

SOURCE: U.S., 19 pp.

CODEN: USXXAM

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

INVENTOR(S):

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6355756	B1	20020312	US 1999-313903	19990518
US 2001024738	A1	20010927	US 2001-819119	20010327
US 6433115	B2	20020813		
US 2002045719	A1	20020418	US 2001-819116	20010327
US 6512070	B2	20030128		
PRIORITY APPLN. INFO.:			US 1999-313903	A3 19990518

ED Entered STN: 14 Mar 2002

AB A process for preparing a dual purpose electroactive copolymer comprised of charge transport polymeric segments and a light emitting polymeric segment comprises: (a) contacting a dihalo-substituted polycyclic aromatic reactant with a living free radical polymerization initiator under conditions effective to bring about polymerization, resulting in a light emitting polymeric intermediate comprised of linked polycyclic aromatic monomer units and two or more displaceable termini; and (b) synthesizing a charge transport polymeric segment comprised of polymerized charge transporting monomer units at each of the displaceable termini, via living free radical polymerization The copolymers may be A-B-A triblock copolymers, brush-type graft copolymers, or variations thereof.

402829-52-1P ΤТ

> (dual purpose electroactive copolymers, preparation thereof, and use in opto-electronic devices)

> > PAGE 1-A

RN 402829-52-1 HCAPLUS

CN Poly(9,9-dihexyl-9H-fluorene-2,7-diyl),  $\alpha$ , $\omega$ -bis[4-[1-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl]phenyl]- (9CI) INDEX NAME)

Me Me— (CH2)5 M⊖  $(CH_2)_5$  — Me M℮ ςн-

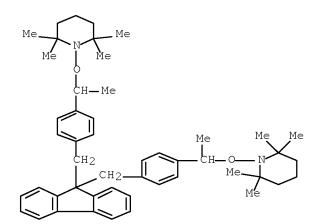
PAGE 1-B

IT 402829-51-0P

(dual purpose electroactive copolymers, preparation thereof, and use in opto-electronic devices)

RN 402829-51-0 HCAPLUS

CN Piperidine, 1,1'-[9H-fluoren-9-ylidenebis(methylene-4,1-phenyleneethylideneoxy)]bis[2,2,6,6-tetramethyl-(9CI) (CA INDEX NAME)



IC ICM C08F112-06

INCL 526347100

CC 35-4 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 73

IT Polymerization catalysts

(living, radical; dual purpose electroactive copolymers, preparation thereof, and use in opto-electronic devices)

IT 9003-53-6DP, Polystyrene, reaction products with fluorenyl piperidine derivs. 25067-59-8DP, N-Vinylcarbazole homopolymer, reaction products with piperidinyloxy compds. 201807-75-2DP,

TEMPO-functionalized 225936-03-8P 362524-25-2DP, TEMPO-functionalized 402829-49-6P 402829-50-9P

402829-52-1P

(dual purpose electroactive copolymers, preparation thereof, and use in opto-electronic devices)

IT 236092-91-4P 402829-51-0P

(dual purpose electroactive copolymers, preparation thereof, and use in opto-electronic devices)

REFERENCE COUNT: 25 THERE ARE 25 CITED REFERENCES AVAILABLE FOR

THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L69 ANSWER 21 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2001:629050 HCAPLUS Full-text

DOCUMENT NUMBER: 135:358194

TITLE: Synthesis of three- and six-arms polystyrene via

living/controlled free radical polymerization

AUTHOR(S): Chessa, G.; Scrivanti, A.; Matteoli, U.;

Castelvetro, V.

CORPORATE SOURCE: Dipartimento di Chimica, Universita di Venezia,

Venice, 30123, Italy

SOURCE: Polymer (2001), 42(23), 9347-9353 CODEN: POLMAG; ISSN: 0032-3861

PUBLISHER: Elsevier Science Ltd.

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 30 Aug 2001

Three new polyfunctional TEMPO-based initiators have been synthesized by reaction of some mesitylene cores and 2,2,6,6-tetramethyl-1-(2-hydroxy-1-phenylethoxy)-piperidine. They have been employed in the living/controlled radical polymerization of styrene to provide three- and six-arm star macromols. These polymers have mol. weight ranging from 6000 to 11000 g/mol and narrow mol. weight distributions (PD<1.3). Cleavage of the link between the core and the arms was achieved, using a two-step sequence implying the preliminary removal of the TEMPO chain ends followed by catalytic hydrogenolysis. The dimensions of the individual arms so obtained closely match the values expected from the styrene/initiator molar ratio in the polymerization feed.

IT 372522-45-7P 372522-46-8P 372522-47-9P

(synthesis of three- and six-arm polystyrene via living/controlled free radical polymerization)

RN 372522-45-7 HCAPLUS

CN Piperidine, 1,1',1''-[1,3,5-benzenetriyltris[methyleneoxy(1-phenyl-2,1-ethanediyl)oxy]]tris[2,2,6,6-tetramethyl- (9CI) (CA INDEX NAME)

RN 372522-46-8 HCAPLUS

CN Piperidine, 1,1',1''-[(2,4,6-trimethyl-1,3,5-benzenetriyl)tris[methyleneoxy(1-phenyl-2,1-

ethanediyl)oxy]]tris[2,2,6,6-tetramethyl- (9CI) (CA INDEX NAME)

RN 372522-47-9 HCAPLUS

CN Pyridine, 4,4',4''-[1,3,5-benzenetriyltris(methyleneoxy)]tris[2,6-bis[[2-phenyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethoxy]methyl]-(9CI) (CA INDEX NAME)

PAGE 1-B

PAGE 2-B

- CC 35-3 (Chemistry of Synthetic High Polymers)
- ST polystyrene multiarm living controlled radical polymn; TEMPO based catalyst multiarm polystyrene; star branched polystyrene TEMPO deriv catalyst
- IT Polymerization catalysts

(synthesis of three- and six-arm polystyrene via living/controlled free radical polymerization)

IT 18226-42-1, 1,3,5-Tris(bromomethyl)benzene 161776-41-6 (in catalyst preparation; synthesis of three- and six-arm

polystyrene via living/controlled free radical polymerization)

IT 372522-45-7P 372522-46-8P 372522-47-9P

(synthesis of three- and six-arm polystyrene via living/controlled free radical polymerization)

REFERENCE COUNT:

THERE ARE 25 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L69 ANSWER 22 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2001:208762 HCAPLUS Full-text

25

DOCUMENT NUMBER: 135:5896

TITLE: Synthesis of poly(methylene-b-styrene) by

sequential living polymerization Zhou, Xian-Zhi; Shea, Kenneth J.

AUTHOR(S): Zhou, Xian-Zhi; Shea, Kenneth J. CORPORATE SOURCE: Department of Chemistry, University of California

Irvine, Irvine, CA, 92697-2025, USA

SOURCE: Macromolecules (2001), 34(9), 3111-3114

CODEN: MAMOBX; ISSN: 0024-9297

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 23 Mar 2001

AB Poly homologation reaction for the synthesis of poly(methylene-b- styrene) block copolymers was considered. A series of block copolymers were prepared by the hydroboration-poly homologation. Control over the chain length of the polymethylene block was achieved by adjusting the initial molar ratio of ylide to organoborane.

IT 341968-37-4P

(borane; synthesis of poly(methylene-b-styrene) by sequential living polymerization)

RN 341968-37-4 HCAPLUS

CN Piperidine, 1,1',1''-[borylidynetris[3,1-propanediyloxy(1-phenyl-2,1-ethanediyl)oxy]]tris[2,2,6,6-tetramethyl- (9CI) (CA INDEX NAME)

CC 35-4 (Chemistry of Synthetic High Polymers)

IT 341968-37-4P

(borane; synthesis of poly(methylene-b-styrene) by sequential living polymerization)

IT 161776-41-6P

(initiator; synthesis of poly(methylene-b-styrene) by sequential living polymerization)

REFERENCE COUNT: 22 THERE ARE 22 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

RE FORM

L69 ANSWER 23 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 2001:31452 HCAPLUS Full-text DOCUMENT NUMBER: 134:101275

TITLE: Preparation of mono and multifunctional alkoxyamines for forming nitroxyl radical

initiators and regulators useful in the

preparation of polymers with narrow polydispersity

INVENTOR(S): Kramer, Andreas; Nesvadba, Peter; Zink,

Marie-Odile; Wunderlich, Wiebke

PATENT ASSIGNEE(S): Ciba Specialty Chemicals Holding Inc., Switz.

SOURCE: PCT Int. Appl., 74 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

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	2001002345 2001002345						2001 2001			WO 2	2000-1	EP58	99		2	0000	626
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		CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EE,	ES,	FΙ,	GB,	GD,	GE,	GH,	GM,	
		HR,	HU,	ID,	IL,	IN,	IS,	JP,	KΕ,	KG,	KP,	KR,	KΖ,	LC,	LK,	LR,	
		LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	${ m MZ}$ ,	NO,	NΖ,	PL,	
		PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	ТJ,	TM,	TR,	TT,	TZ,	UA,	
		UG,	US,	UZ,	VN,	YU,	ZA,	ZW,	AM,	ΑZ,	BY,	KG,	KΖ,	MD,	RU,	ТJ,	TM
	RW:	GH,	GM,	ΚE,	LS,	MW,	MZ,	SD,	SL,	SZ,	TZ,	UG,	ZW,	ΑT,	BE,	CH,	
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		BF,	ВJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GW,	ML,	MR,	ΝE,	SN,	TD,	ΤG	
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EP	EP 1189875					A2 20020327			EP 2000-951302						20000626		
EP	1189	875			В1		2004	0804									
	R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	ΙΤ,	LI,	LU,	NL,	MC,	PT,	
		ΙE,	SI,	LT,	LV,	FI,	RO										
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PRIORIT	Y APP	LN.	INFO	.:						EP 1	.999-	8105	67		A 1	9990	702
										WO 2	2000-1	EP58	99	1	W 2	0000	626

OTHER SOURCE(S): MARPAT 134:101275

ED Entered STN: 12 Jan 2001

AB The title alkoxyamines especially useful for the living polymerization of unsatd. monomers or/and oligomers giving polymers with good conversion are compds. bearing groups which can liberate stable free nitroxyl radicals of specific structures.

IT 319457-98-2P 319457-99-3P 319458-00-9P 319458-01-0P 319458-02-1P 319458-05-4P 319458-06-5P 319458-07-6P 319458-09-8P 319458-10-1P 319458-13-4P 319458-14-5P 319458-18-9P 319458-19-0P 319458-20-3P 319458-21-4P 319458-22-5P 319458-23-6P 319458-24-7P 319458-27-0P 319458-29-2P 319458-32-7P 319458-34-9P 319458-37-2P

319458-40-7P 319458-43-0P 319458-46-3P 319458-49-6P 319458-51-0P 319458-54-3P

(initiator; preparation of mono and multifunctional alkoxyamines as initiators for free radical polymerization with narrow polydispersity)

RN 319457-98-2 HCAPLUS

CN 4-Piperidinol, 1,1',1''-[1,3,5-triazine-2,4,6-triyltris(oxy-4,1-phenyleneethylideneoxy)]tris[2,2,6,6-tetramethyl- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 2-B

\_\_\_OH

RN 319457-99-3 HCAPLUS

CN 4-Piperidinol, 1,1'-[1,4-phenylenebis(methyleneoxy-4,1-phenyleneethylideneoxy)]bis[2,2,6,6-tetramethyl-(9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 319458-00-9 HCAPLUS

CN 4-Piperidinol, 1,1'-[[1,1'-biphenyl]-4,4'-diylbis(methyleneoxy-4,1-phenyleneethylideneoxy)]bis[2,2,6,6-tetramethyl- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 319458-01-0 HCAPLUS

CN 4-Piperidinol, 1,1',1'',1'''-[1,2,4,5-benzenetetrayltetrakis(methylene oxy-4,1-phenyleneethylideneoxy)]tetrakis[2,2,6,6-tetramethyl- (9CI) (CA INDEX NAME)

PAGE 1-B

RN 319458-02-1 HCAPLUS
CN 4-Piperidinol, 1,1',1'',1''',1''''-[1,2,3,4,5,6-benzenehexaylhexakis(methyleneoxy-4,1-phenyleneethylideneoxy)]hexakis[2,2,6,6-tetramethyl- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

#### PAGE 2-B

PAGE 3-A
Me
OH

RN 319458-05-4 HCAPLUS

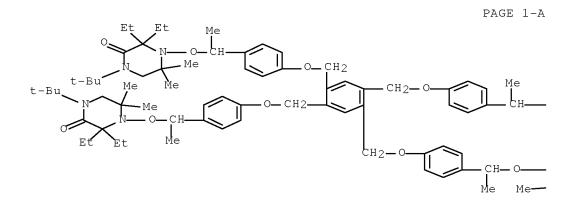
CN Piperazinone, 4,4'-[1,4-phenylenebis(methyleneoxy-4,1-phenyleneethylideneoxy)]bis[1-(1,1-dimethylethyl)-3,3-diethyl-5,5-dimethyl- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 319458-06-5 HCAPLUS

CN Piperazinone, 4,4',4'',4'''-[1,2,4,5-benzenetetrayltetrakis(methyleneo xy-4,1-phenyleneethylideneoxy)]tetrakis[1-(1,1-dimethylethyl)-3,3-diethyl-5,5-dimethyl- (9CI) (CA INDEX NAME)



RN 319458-07-6 HCAPLUS

CN Piperazinone, 4,4',4'',4''',4'''',4''''-[1,2,3,4,5,6-benzenehexaylhexakis(methyleneoxy-4,1-phenyleneethylideneoxy)]hexakis[1-(1,1-dimethylethyl)-3,3-diethyl-5,5-dimethyl-(9CI) (CA INDEX NAME)

PAGE 2-B

RN 319458-09-8 HCAPLUS

CN Benzoic acid, 4-[1-[[4-(1,1-dimethylethyl)-2,2-diethyl-6,6-dimethyl-3-oxo-1-piperazinyl]oxy]ethyl]-, 1,12-dodecanediyl ester (9CI) (CA INDEX NAME)

PAGE 1-B

RN 319458-10-1 HCAPLUS

CN Benzoic acid, 4-[1-[[4-(1,1-dimethylethyl)-2,2-diethyl-6,6-dimethyl-3-oxo-1-piperazinyl]oxy]ethyl]-, 2,2-bis[[[4-[1-[[4-(1,1-dimethylethyl)-2,2-diethyl-6,6-dimethyl-3-oxo-1-piperazinyl]oxy]ethyl]benzoyl]oxy]methyl]-1,3-propanediyl ester (9CI) (CA INDEX NAME)

# PAGE 1-A

# PAGE 2-B

$$Me \xrightarrow{N} Et$$

$$t-Bu$$

RN 319458-13-4 HCAPLUS

CN 4-Piperidinol, 1,1'-[1,4-phenylenebis(methyleneoxy-4,1-phenyleneethylideneoxy)]bis[2,6-diethyl-2,3,6-trimethyl-(9CI) (CA INDEX NAME)

### PAGE 1-B

RN 319458-14-5 HCAPLUS

CN 4-Piperidinol, 1,1',1'',1'''-[1,2,4,5-benzenetetrayltetrakis(methylene oxy-4,1-phenyleneethylideneoxy)]tetrakis[2,6-diethyl-2,3,6-trimethyl-(9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 319458-18-9 HCAPLUS

CN 1,4-Benzenedicarboxylic acid, bis[1-[4-[1-[[4-(benzoyloxy)-2,2,6,6-tetramethyl-1-piperidinyl]oxy]ethyl]phenyl]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

$$\begin{array}{c} \text{Me} & \text{Me} \\ \text{Me} & \text{Me} \\ \text{N} & \text{O-CH} \\ \text{Me} \end{array}$$

PAGE 1-B

RN 319458-19-0 HCAPLUS

CN Ethanedioic acid, bis[1-[4-[1-[[4-(benzoyloxy)-2,2,6,6-tetramethyl-1-piperidinyl]oxy]ethyl]phenyl]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 319458-20-3 HCAPLUS

CN Octanedioic acid, bis[1-[4-[1-[[4-(benzoyloxy)-2,2,6,6-tetramethyl-1-piperidinyl]oxy]ethyl]phenyl]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 319458-21-4 HCAPLUS

CN Decanedioic acid, bis[1-[4-[1-[[4-(benzoyloxy)-2,2,6,6-tetramethyl-1-piperidinyl]oxy]ethyl]phenyl]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 319458-22-5 HCAPLUS

CN 4-Piperidinol, 1,1'-[1,4-phenylenebis(methyleneoxyethylidene-4,1-phenyleneethylideneoxy)]bis[2,2,6,6-tetramethyl-, dibenzoate (ester) (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 319458-23-6 HCAPLUS

CN Carbamic acid, 1,6-hexanediylbis-, bis[1-[4-[1-[[4-(benzoyloxy)-2,2,6,6-tetramethyl-1-piperidinyl]oxy]ethyl]phenyl]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 319458-24-7 HCAPLUS

CN 4-Piperidinol, 1,1',1''-[1,3,5-triazine-2,4,6-triyltris(oxyethylidene-4,1-phenyleneethylideneoxy)]tris[2,2,6,6-tetramethyl-, tribenzoate (ester) (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 2-B

RN 319458-27-0 HCAPLUS

CN Octanedioic acid, bis[1-[4-[1-[[2,2,6,6-tetramethyl-4-[(2,4,6-trimethylbenzoyl)oxy]-1-piperidinyl]oxy]ethyl]phenyl]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 319458-29-2 HCAPLUS

CN Octanedioic acid, bis[1-[4-[1-[[4-(2,2-dimethyl-1-oxopropoxy)-2,2,6,6-tetramethyl-1-piperidinyl]oxy]ethyl]phenyl]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 319458-32-7 HCAPLUS

CN Octanedioic acid, bis[1-[4-[1-[[2,2,6,6-tetramethyl-4[(trimethylsilyl)oxy]-1-piperidinyl]oxy]ethyl]phenyl]ethyl] ester
(9CI) (CA INDEX NAME)

PAGE 1-B

RN 319458-34-9 HCAPLUS

CN Octanedioic acid, bis[1-[4-[1-[(2,2,6,6-tetramethyl-4-propoxy-1-piperidinyl)oxy]ethyl]phenyl]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-B

RN 319458-37-2 HCAPLUS

CN Octanedioic acid, bis[1-[4-[1-[[4-(benzoyloxy)-2,2-diethyl-6,6-dimethyl-1-piperidinyl]oxy]ethyl]phenyl]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 319458-40-7 HCAPLUS

CN Octanedioic acid, bis[1-[4-[1-[[4-(benzoyloxy)-2,6-diethyl-2,3,6-trimethyl-1-piperidinyl]oxy]ethyl]phenyl]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

$$\begin{array}{c} \text{Me} \\ \text{Me} \\ \text{N} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{Me} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \end{array} \begin{array}{c} \text{O} \\ \text{CH} \\ \text{O} \\ \text{CH} \\ \text{O} \\ \text{O} \\ \text{CH} \\ \text{O} \\ \text{O} \\ \text{O} \\ \text{O} \\ \text{CH} \\ \text{O} \\ \text{$$

PAGE 1-B

RN 319458-43-0 HCAPLUS

CN Octanedioic acid, bis[1-[4-[1-[[4-(acetylamino)-2,2,6,6-tetramethyl-1-piperidinyl]oxy]ethyl]phenyl]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-B

RN 319458-46-3 HCAPLUS

CN Octanedioic acid, bis[1-[4-[1-[[4-(benzoylamino)-2,6-diethyl-2,3,6-trimethyl-1-piperidinyl]oxy]ethyl]phenyl]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 319458-49-6 HCAPLUS

CN Octanedioic acid, bis[1-[4-[1-[(2,6-diethyl-2,3,6-trimethyl-4-oxo-1-piperidinyl)oxy]ethyl]phenyl]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-B

RN 319458-51-0 HCAPLUS

CN Octanedioic acid, bis[1-[4-[1-[[4-(1,1-dimethylethyl)-2,2-diethyl-6,6-dimethyl-3-oxo-1-piperazinyl]oxy]ethyl]phenyl]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-B

RN 319458-54-3 HCAPLUS

CN Octanedioic acid, bis[1-[4-[3,4-bis(1,1-dimethylethyl)-5-ethoxy-1-methyl-5-oxido-2,6-dioxa-3-aza-5-phosphaoct-1-yl]phenyl]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

- IC ICM C07C239-20
  - ICS C07D211-94; C08F004-00
- CC 35-3 (Chemistry of Synthetic High Polymers)
- ST alkoxyamine multifunctional nitroxyl initiator radical living polymn; narrow polydispersity radical living polymn; mol wt distribution living polymn nitroxyl radical initiator

```
Amines, preparation
ΙT
        (alkoxylated, derivs., initiator; preparation of mono and
       multifunctional alkoxyamines as initiators for free
       radical polymerization with narrow polydispersity)
ΙT
    Polymerization
        (living, radical; preparation of mono and multifunctional alkoxyamines
        as initiators for free radical polymerization with narrow
       polydispersity)
TΤ
    Polymerization catalysts
        (living; preparation of mono and multifunctional alkoxyamines as
        initiators for free radical polymerization with narrow
       polydispersity)
    Nitroxides
ΙT
        (preparation of mono and multifunctional alkoxyamines as
        initiators for free radical polymerization with narrow
       polydispersity)
ΙT
    Polymerization catalysts
        (radical; preparation of mono and multifunctional alkoxyamines as
        initiators for free radical polymerization with narrow
       polydispersity)
    243972-13-6P 243972-14-7P 243972-16-9P 264280-52-6P
ΙT
    319457-95-9P 319457-96-0P 319457-97-1P 319458-04-3P
    319458-08-7P 319458-11-2P 319458-12-3P 319458-15-6P
    319458-16-7P 319458-17-8P 319458-25-8P 319458-26-9P
    319458-28-1P 319458-30-5P 319458-31-6P 319458-33-8P
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                                                 319458-39-4P
                                  319458-44-1P
    319458-41-8P 319458-42-9P
                                                 319458-45-2P
    319458-47-4P 319458-48-5P 319458-50-9P 319458-52-1P
    319458-53-2P
        (initiator/intermediate for multifunctional
       initiator; preparation of mono and multifunctional alkoxyamines
        as initiators for free radical polymerization with narrow
       polydispersity)
ΙT
    264280-51-5P 319457-98-2P 319457-99-3P
    319458-00-9P 319458-01-0P 319458-02-1P
    319458-03-2P 319458-05-4P 319458-06-5P
    319458-07-6P 319458-09-8P 319458-10-1P
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    319458-22-5P 319458-23-6P 319458-24-7P
    319458-27-0P 319458-29-2P 319458-32-7P
    319458-34-9P 319458-37-2P 319458-40-7P
    319458-43-0P 319458-46-3P 319458-49-6P
    319458-51-0P 319458-54-3P
        (initiator; preparation of mono and multifunctional
        alkoxyamines as initiations for free radical polymerization with
        narrow polydispersity)
    9003-49-0P, Butyl acrylate homopolymer
ΙT
                                             9003-53-6P, Polystyrene
        (linear or star-shaped; preparation of mono and multifunctional
        alkoxyamines as initiators for free radical polymerization with
        narrow polydispersity)
    106-95-6, Allyl bromide, reactions 108-77-0, Cyanuric chloride
ΤТ
    115-77-5, Pentaerythritol, reactions
                                          535-11-5, Ethyl
    2-bromopropionate 623-24-5, \alpha, \alpha'-Dibromo-p-xylene
    814-68-6, Acryloyl chloride 937-30-4 2226-96-2,
    4-Hydroxy-2,2,6,6-tetramethylpiperidin-1-oxyl
                                                    3095-73-6,
    Hexakis(bromomethyl)benzene 3245-23-6, 4-Ethylphenyl acetate
    5675-51-4, 1,12-Dodecanediol 15442-91-8, 1,2,4,5-
    Tetrakis (bromomethyl) benzene 20248-86-6, 4,4'-
    Bis (bromomethyl) biphenyl 61745-37-7 264279-93-8
```

(reactant for initiator; preparation of mono and multifunctional alkoxyamines as initiators for free radical polymerization with narrow polydispersity)

L69 ANSWER 24 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1999:279746 HCAPLUS Full-text

DOCUMENT NUMBER: 130:325501

TITLE: Procedure for manufacture of block copolymers with

controlled architecture via functional radical

initiators and living radical polymerization, and initiator

compositions, and corresponding copolymers

INVENTOR(S):
Bertin, Denis; Destarac, Mathias; Boutevin,

Bernard

PATENT ASSIGNEE(S): Elf Atochem S.A., Fr. SOURCE: Eur. Pat. Appl., 34 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent LANGUAGE: French

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PA	PATENT NO.						DATE		APPLICATION NO.	DATE		
EP	911350			A1 1		19990428		EP 1998-402624	19981022			
	R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB, GR, IT, LI, LU, NL, S	E, MC,		
		PT,	ΙE,	SI,	LT,	LV,	FΙ,	RO				
FR	2770	219			A1		1999	0430	FR 1997-13383	19971024		
EP	1277	771			A2		2003	0122	EP 2002-78512	19981022		
	R:	ΑT,	BE,	DE,	ES,	FR,	GB,	ΙΤ,	NL, SE, FI			
EP	1288	236			A2		2003	0305	EP 2002-78511	19981022		
	R:	ΑT,	BE,	DE,	ES,	FR,	GB,	ΙΤ,	NL, SE, FI			
PRIORIT	Y APP	LN.	INFO	.:					FR 1997-13383 A	19971024		
									EP 1998-402624 A3	19981022		

OTHER SOURCE(S): MARPAT 130:325501

ED Entered STN: 06 May 1999

The first step in the procedure is thermally induced radical polymerization of AΒ at least one monomer M1 = methacrylic monomer and an initiator X-A-Y, where X and Y = monovalent living radical groups and A = divalent linking group capable of radical polymerization and radical-reactive groups, e.g., diazo -N=N-, peroxide -O-O-, to obtain a living polymer X-D (or Y-E)PM1-T where D = is a free radical moiety and T = terminal group. The initiator is obtained by reaction of 4,4'-Azobis[4-cyanovaleric acid] and end-functionalized alcs. or by reaction of H2O2 and an acid chloride containing groups X or Y. In the second step, living radical polymerization of the polymer and at least one other monomer, M2, is carried out, either by photochem. activation or by chain transfer control, to obtain a multifunctional macroinitiator that can be used in yet another polymerization step with at least one monomer M3; M2 and M3 are selected from vinyl, allyl, vinylidene, diene, or olefinic monomers. The second step is carried out in presence of transition metal complex catalysts , preferably CuZ'/L where Z = halogen, hexafluorophosphate, acetate and L =  $\alpha$ diimine ligand. Thus, a triblock copolymer, PS-PABu-PS was obtained; the initiator was prepared from 4,4'-Azobis[4-cyanovaleric acid] and trichloroethanol and used in radical polymerization of Bu acrylate at  $130\,^{\circ}$  to obtain the living poly(Bu acrylate) of average mol. weight 8700 g/mol and degree of polymerization of 64. The living polymer was then mixed with styrene and CuCl and bipyridine as radical polymerization catalyst system; the triblock copolymer was isolated from the reaction medium and has average mol.

weight of 92,600 g/mol, of which 8700 g/mol correspond to the poly(Bu acrylate) sequence and 45,000 to the polystyrene sequences.

IT 223668-08-4P

(peroxide initiator; functional radical

initiators in sequential radical and living radical polymerization for manufacture of block copolymers with controlled architecture)

RN 223668-08-4 HCAPLUS

CN Benzoic acid, 4,4'-(dioxydicarbonyl)bis-, bis[2-phenyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

Me Me Ph
N O CH CH2 O CH2

PAGE 1-B

IC ICM C08F293-00

ICS C07C255-65

CC 35-4 (Chemistry of Synthetic High Polymers)

IT Polymerization

Polymerization

(block, photochem.; functional radical initiators in sequential radical and living radical polymerization for manufacture of

copolymers with controlled architecture)

IT Polymerization

block

block

Polymerization catalysts

(living; functional radical initiators in sequential

 $\,$  radical and living radical polymerization for manufacture of block copolymers

with controlled architecture)

IT Polymerization catalysts

(photopolymn., block; functional radical initiators in sequential radical and living radical polymerization for manufacture of

copolymers with controlled architecture)

IT Polymerization

Polymerization catalysts

(radical; functional radical initiators in sequential

radical and living radical polymerization for manufacture of block copolymers

with controlled architecture)

IT 213546-64-6P 223668-06-2P

(azobis initiator; functional radical initiators in sequential radical and living radical polymerization for manufacture of block copolymers with controlled architecture) ΤТ 106399-43-3P, Butadiene-methyl methacrylate block copolymer 725713-28-0P (diblock; functional radical initiators in sequential radical and living radical polymerization for manufacture of block copolymers with controlled architecture) 100-20-9, 1,4-Benzenedicarbonyl dichloride 115-20-8, Trichloroethanol 2638-94-0, 4,4'-Azobis[4-cyanovaleric acid] 7722-84-1, Hydrogen peroxide (H2O2), reactions 30887-99-1 161776-41-6, 2-Phenyl-2-[(2,2,6,6-tetramethylpiperidino)oxy]ethanol(functional radical initiators in sequential radical and living radical polymerization for manufacture of block copolymers with controlled architecture) ΙT 9003-49-0P, Poly(butyl acrylate) 9011-14-7P, Poly(methyl methacrylate) (functional radical initiators in sequential radical and living radical polymerization for manufacture of block copolymers with controlled architecture) 223668-07-3P ΤТ (intermediate; functional radical initiators in sequential radical and living radical polymerization for manufacture of block copolymers with controlled architecture) 223668-08-4P TT (peroxide initiator; functional radical initiators in sequential radical and living radical polymerization for manufacture of block copolymers with controlled architecture) 7758-89-6, Copper chloride (CuCl) 37275-48-2, Bipyridine ΙT (radical polymerization catalyst system; functional radical initiators in sequential radical and living radical polymerization for manufacture of block copolymers with controlled architecture) 110772-34-4P, Butyl acrylate-styrene block copolymer 705279-67-0P ΤТ (triblock; functional radical initiators in sequential radical and living radical polymerization for manufacture of block copolymers with controlled architecture) THERE ARE 7 CITED REFERENCES AVAILABLE FOR REFERENCE COUNT: 7 THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L69 ANSWER 25 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1998:572917 HCAPLUS Full-text DOCUMENT NUMBER: 129:276766 TITLE: Soluble supports tailored for organic synthesis: parallel polymer synthesis via sequential normal/living free radical processes AUTHOR(S): Gravert, Dennis J.; Datta, Anita; Wentworth, Paul, Jr.; Janda, Kim D. CORPORATE SOURCE: Department of Chemistry and The Skaggs Institute for Chemical Biology, Scripps Research Institute, La Jolla, CA, 92037, USA SOURCE: Journal of the American Chemical Society (1998), 120(37), 9481-9495 CODEN: JACSAT; ISSN: 0002-7863 PUBLISHER: American Chemical Society DOCUMENT TYPE: Journal English LANGUAGE:

ED Entered STN: 10 Sep 1998

AΒ To expand the availability and solubility range of polymer supports for liquid-phase organic synthesis (LPOS) we have applied a sequence of normal and "living" free radical polymerization to generate a library of block copolymers possessing either block or graft architecture with imitiators and a diverse set of vinyl monomers. The structure, mol. weight, and polydispersity (PD) of the individual library members have been determined by size exclusion chromatog. (SEC), 1H and 13C NMR, and as a function of the solubility of each polymer in a range of solvents. One copolymer, polyBS-DS (Mn = 17 000, PD = 1.54) derived from 4-tert-butylstyrene (BS), 3,4-dimethoxystyrene (DS) has a solubility profile [soluble in toluene, THF (THF), ether, acetone and methylene chloride (DCM), insol. in methanol and water] that is different from the present polymer of choice for LPOS, poly(ethylene) glycol (PEG), and was studied in some detail as a new support in LPOS. The lpha-nitrile groups of polyBS-DS are reduced smoothly with LiAlH4 in THF to give the amino functionalized copolymer (0.14 mmol g-1 of amino groups based on a quant. ninhydrin anal.). Kinetic studies have revealed that derivatization of the amino groups of the copolymer with 4-dimethylaminocinnamaldehyde occurs at a comparable rate to a solution counterpart (kpoly22 = 0.49 L mol-1 h-1 vskaminohexane = 0.69 L mol-1 h-1). Following reaction with N-glutaroyl-(2S, 4S)-4-diphenylphosphino- 2-[(diphenylphosphino)methyl]pyrrolidine and exchange of Rh(I), the resulting phosphine containing copolymer, catalyzes the enantioselective hydrogenation of 2-N-acetamidoacrylic acid to N-acetylalanine in THF. An 87% enantiomeric excess (ee) of (S)-N-acetylalanine is obtained, comparable to that observed with a homogeneous phosphine ligand. This work highlights the power of a parallel polymer synthesis strategy, from conception to application, for the generation of polymers possessing unique solubility profiles and functionality which can serve as novel supports in LPOS.

IT 188119-33-7 203382-60-9

(catalyst for; parallel polymer preparation via sequential normal/living free radical polymerization)

RN 188119-33-7 HCAPLUS

CN Pentanoic acid, 4,4'-azobis[4-cyano-, bis[2-phenyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-B

RN 203382-60-9 HCAPLUS

CN Pentanenitrile, 2,2'-azobis[2-methyl-5-[2-phenyl-2-[(2,2,6,6-

tetramethyl-1-piperidinyl)oxy]ethoxy]- (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

ΙT 213994-38-8P

> (catalyst for; parallel polymer preparation via sequential normal/living free radical polymerization)

RN 213994-38-8 HCAPLUS

Pentanoic acid, 4,4'-azobis[4-cyano-, bis[2-[[4-[[(1,1-CN dimethylethoxy)carbonyl]amino]-2,2,6,6-tetramethyl-1-piperidinyl]oxy]-2-phenylethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

$$t-BuO-\stackrel{\bigcirc}{C}-NH \xrightarrow{Me} \stackrel{Me}{\longrightarrow} \stackrel{Me}$$

PAGE 1-B

$$-CH_2 - CH_2 -$$

CC 37-3 (Plastics Manufacture and Processing) Section cross-reference(s): 67

dimethoxystyrene butylstyrene block polymer datalyst support ST

ΙT 188119-33-7 203382-60-9

(catalyst for; parallel polymer preparation via sequential normal/living free radical polymerization) ΙT 213994-38-8P (catalyst for; parallel polymer preparation via sequential normal/living free radical polymerization) ΙT 920-46-7, Methacryloyl chloride 4693-47-4 14691-88-4 24424-99-5, Di-tert-butyl dicarbonate 161776-41-6 (in preparation of catalyst; parallel polymer preparation via sequential normal/living free radical polymerization) 213994-43-5P 213994-47-9P 213994-50-4P ΙT (in preparation of catalyst; parallel polymer preparation via sequential normal/living free radical polymerization) 12092-47-6,  $\mu$ -Dichloro-bis(1,5-cyclooctadiene)-dirhodium ΤT 213994-93-5D, reaction products with reduced and hydrogenated Bu styrene-dimethoxystyrene block copolymer (in sequential normal/living free radical polymerization for polymer-supported catalyst) 61478-29-3 ΙT (in sequential normal/living free radical polymerization for polymer-supported catalyst) ΙT 213994-64-0DP, reduced with LiAlH4, hydrogenated, derivs. (sequential normal/living free radical polymerization for polymer-supported catalyst) ΙT 5429-56-1 (sequential normal/living free radical polymerization for polymer-supported catalyst for hydrogenation) ΙT 97-69-8P (sequential normal/living free radical polymerization for polymer-supported catalyst for hydrogenation) REFERENCE COUNT: 85 THERE ARE 85 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE L69 ANSWER 26 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1997:532628 HCAPLUS Full-text DOCUMENT NUMBER: 127:221034 TITLE: Block Copolymer Preparation Using Sequential Normal/Living Radical Polymerization Techniques Li, I. Q.; Howell, B. A.; Dineen, M. T.; Kastl, P. AUTHOR(S): E.; Lyons, J. W.; Meunier, D. M.; Smith, P. B.; Priddy, D. B. Center for Applications in Polymer Science, CORPORATE SOURCE: Central Michigan University, Mount Pleasant, MI, 48859, USA SOURCE: Macromolecules (1997), 30(18), 5195-5199 CODEN: MAMOBX; ISSN: 0024-9297 American Chemical Society PUBLISHER: DOCUMENT TYPE: Journal LANGUAGE: English Entered STN: 21 Aug 1997 EDAnionic and nitroxide-mediated (NM) radical polymerization works well for AB styrene but not for acrylates. We have overcome this problem and successfully prepared styrene-b-Bu acrylate (S-BA), styrene-b-Me methacrylate (S-MMA), styrene-b-isoprene (S-IP), and styrene-alt-acrylonitrile-b-isoprene (SAN-IP) polymers using a sequential normal/living radical polymerization scheme. Clear (S-IP and SAN-IP) to translucent (S-BA and S-MMA) films were obtained having microphase-separated polymer morphol. GPC studies and chemical digestion of the IP segments of S-IP and SAN-IP block copolymers confirmed their block structure. The sequential normal/living radical polymerization

approach provides a new route to synthesize block polymers that have previously proven very difficult to make.

IT 188119-33-7E

(preparation of acrylic block copolymers using sequential normal/living radical polymerization initiated by AIBN and)

RN 188119-33-7 HCAPLUS

CN Pentanoic acid, 4,4'-azobis[4-cyano-, bis[2-phenyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

CC 35-4 (Chemistry of Synthetic High Polymers)
 Section cross-reference(s): 36

ST normal living sequential polymn acrylate methacrylate; isoprene acrylate styrene block copolymer prepn; nitroxide functional azo initiator acrylate polymn; translucent microphase sepd block copolymer synthesis; acrylonitrile isoprene styrene alternating block copolymer; SAN isoprene block copolymer prepn; polyacrylate macroinitiator isoprene styrene polymn

IT 17170-81-9 161776-41-6

(initiator synthesis; preparation of acrylic block copolymers using sequential normal/living radical polymerization)

IT 188119-33-7P

(preparation of acrylic block copolymers using sequential normal/living radical polymerization initiated by AIBN and)  $\,$ 

IT 78-67-1, AIBN

(preparation of acrylic block copolymers using sequential normal/living radical polymerization initiated by nitroxide functional azo initiator and)

REFERENCE COUNT:

THERE ARE 43 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L69 ANSWER 27 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1997:324791 HCAPLUS Full-text

DOCUMENT NUMBER: 127:5476

TITLE: Difunctional living free radical polymerization

initiators for vinyl aromatic monomers

INVENTOR(S): Koster, Robert A.; Priddy, Duane B.; Li, Irene

PATENT ASSIGNEE(S): Dow Chemical Co., USA

SOURCE: U.S., 9 pp.
CODEN: USXXAM

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 5627248	A	19970506	US 1995-533799	19950926
US 5677388	A	19971014	US 1996-731216	19961008
PRIORITY APPLN. INFO.:			US 1995-533799	A3 19950926

OTHER SOURCE(S): MARPAT 127:5476

ED Entered STN: 22 May 1997

AB Vinyl aromatic monomers are polymerized in high conversion and low polydispersity using a diffunctional nitroxyl initiator R1[-CR2R3-O-NR4R5]2; R1 = linking group; R2, R3 = H, alkyl, cycloalkyl, activating group, or alkyl bridging group; R4, R5 = alkyl, aryl, or C4-10-alkyl ring.

IT 184646-29-5P 184646-30-8P

(free radical polymerization initiators for vinyl aromatic monomers)

RN 184646-29-5 HCAPLUS

CN Piperidine, 1,1'-[1,2-ethanediylbis(4,1-phenylenemethyleneoxy)]bis[2,2,6,6-tetramethyl- (9CI) (CA INDEX NAME)

RN 184646-30-8 HCAPLUS

CN Piperidine, 1,1'-[1,4-phenylenebis(ethylideneoxy)]bis[2,2,6,6-tetramethyl- (9CI) (CA INDEX NAME)

IC ICM C08F002-00

ICS C08F220-10; C08F012-08; B01J031-06

INCL 526217000

CC 35-3 (Chemistry of Synthetic High Polymers) Section cross-reference(s): 28, 67

ST nitroxyl catalyst vinyl arom monomer; styrene polymn nitroxyl catalyst; piperidinyloxy styrene polymn

catalyst; radical polymn catalyst vinyl monomer

IT Polymerization catalysts

(radical, nitroxyl free radicals; free radical polymerization initiators for vinyl aromatic monomers)

IT 154554-67-3P 184646-29-5P 184646-30-8P

(free radical polymerization initiators for vinyl aromatic monomers)

IT 95-78-3 105-05-5 1633-22-3 2564-83-2

(free radical polymerization initiators for vinyl aromatic monomers)

IT 9003-53-6P 121523-76-0P

(free radical polymerization initiators for vinyl aromatic monomers)

L69 ANSWER 28 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1997:224604 HCAPLUS <u>Full-text</u>

DOCUMENT NUMBER: 126:225616

TITLE: Block copolymer preparation using normal/living

tandem polymerization Li, I. Q.; Howell, B. A.

CORPORATE SOURCE: Cent. Appli. Polym. Sci., Central Michigan Univ.,

nmunt Pleasant, MI, 48859, USA

SOURCE: Polymer Preprints (American Chemical Society,

Division of Polymer Chemistry) (1997), 38(1),

762-763

CODEN: ACPPAY; ISSN: 0032-3934

PUBLISHER: American Chemical Society, Division of Polymer

Chemistry

DOCUMENT TYPE: Journal LANGUAGE: English ED Entered STN: 07 Apr 1997

AB Normal/living sequential polymerization of Bu acrylate and styrene yields translucent films having microphase separated morphol., indicative of block copolymers with incompatible segments. The tandem polymerization approach is based on normal and nitroxyl-mediated living radical polymerization. In contrast, poly(Bu acrylate) (pBA)- polystyrene prepared by polymerizing styrene in the presence of pBA yielded a brittle and opaque film having a morphol. of pBA droplets dispersed in polystyrene and no sign of microphase separation

IT 188119-33-7P

AUTHOR(S):

(radical initiator; preparation and morphol. of Bu
acrylate-styrene block copolymer by normal/living tandem polymerization)

RN 188119-33-7 HCAPLUS

CN Pentanoic acid, 4,4'-azobis[4-cyano-, bis[2-phenyl-2-[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]ethyl] ester (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

CC 35-4 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 36

IT 188119-33-7P

(radical initiator; preparation and morphol. of Bu
acrylate-styrene block copolymer by normal/living tandem polymerization)

L69 ANSWER 29 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN ACCESSION NUMBER: 1996:436560 HCAPLUS Full-text

DOCUMENT NUMBER: 125:115236

TITLE: Photochemical synthesis of TEMPO-capped

initiators for "living" free radical

polymerization

AUTHOR(S): Connolly, Terrence J.; Baldovi, M. V.; Mohtat, N.;

Scaiano, J. C.

CORPORATE SOURCE: Dep. Chemistry, Univ. Ottawa, Ottawa, ON, K1N 6N5,

Can.

SOURCE: Tetrahedron Letters (1996), 37(28), 4919-4922

CODEN: TELEAY; ISSN: 0040-4039

PUBLISHER: Elsevier
DOCUMENT TYPE: Journal
LANGUAGE: English
ED Entered STN: 24 Jul 1996

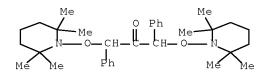
AB Two photochem. routes to stoichiometric initiators used in living free-radical polymns. are presented. These routes offer the advantages of higher yields and allow for the preparation of initiators not accessible using current methodol. All initiators gave detectable carbon centered radicals (laser flash photolysis) and promoted the polymerization of styrene.

IT 179417-97-1P

(catalyst; photochem. synthesis of Tempo-capped initiators for living free-radical polymerization)

RN 179417-97-1 HCAPLUS

CN 2-Propanone, 1,3-diphenyl-1,3-bis[(2,2,6,6-tetramethyl-1-piperidinyl)oxy]- (CA INDEX NAME)



CC 35-3 (Chemistry of Synthetic High Polymers)

ST Tempo capped radical polymn datalyst; piperidinyloxy deriv radical polymn datalyst; styrene radical polymn datalyst

II Polymerization catalysts

```
(living, radical, photochem. synthesis of Tempo-capped free-radical
        initiators for)
ΙT
     179417-96-0P
        (catalyst; photochem. synthesis of TEMPO-capped
        initiators for living free-radical polymerization)
     92952-55-1P
TT
        (catalyst; photochem. synthesis of Tempo-capped
        initiators for living free- radical polymerization)
                  154554-67-3P 157462-14-1P 178625-99-5P
ΙT
     102261-92-7P
     179417-95-9P 179417-97-1P
        (catalyst; photochem. synthesis of Tempo-capped
        initiators for living free-radical polymerization)
     9003-53-6P, Polystyrene
ΙT
        (preparation of; photochem. synthesis of Tempo-capped free-radical
        initiators for)
     70-11-1, Phenacyl bromide 110-05-4, tert-Butyl peroxide
     1,1-Diphenylacetone 958-79-2, 1,3-Dibromo-1,3-diphenyl-2-propanone
     2564-83-2, TEMPO 2632-13-5, 4-Methoxyphenacyl bromide
        (reactant; in photochem. synthesis of Tempo-capped
        initiators for living free-radical polymerization)
L69 ANSWER 30 OF 30 HCAPLUS COPYRIGHT 2008 ACS on STN
ACCESSION NUMBER:
                        1995:734286 HCAPLUS Full-text
DOCUMENT NUMBER:
                        123:144710
TITLE:
                        Architectural control in "living" free radical
                        polymerizations: preparation of star and graft
                        polymers
                        Hawker, Craig J.
AUTHOR(S):
                        Almaden Res. Cent., IBM Res. Cent., San Jose, CA,
CORPORATE SOURCE:
                        95120-6099, USA
SOURCE:
                        Angewandte Chemie, International Edition in
                        English (1995), 34(13/14), 1456-9
                        CODEN: ACIEAY; ISSN: 0570-0833
PUBLISHER:
                        VCH
                         Journal
DOCUMENT TYPE:
LANGUAGE:
                        English
ΕD
     Entered STN: 12 Aug 1995
     Living free radical polymns. based on TEMPO [2,2,6,6-
AΒ
     tetramethylpiperidinyloxy] derivs. alloy for accurate control of macromol.
     architecture. Star and graft copolymers can be prepared from the appropriate
     multi-functional initiators with no crosslinking or termination by
     combination, even under melt conditions. The mol. weight of the arms, or
     grafts, can be controlled by varying the equivalent of monomer added while
     maintaining very low polydispersity. The 2,2,6,6-tetramethylpiperidinyloxy-
     benzoate precursor underwent hydrolysis of the benzyl ester group to give the
     alc. Reaction of the alc. with 1,3,5-benzenetricarbonyl chloride in the
     presence of 4-dimethylaminopyridine produced the tri-functional initiator.
     Bulk polymerization of deuterated styrene with the tri-functional initiator
     produced the polystyrene in 84% yield. The polystyrene underwent hydrolysis
     with KOH and the hydrolyzed product has a mol. weight of 7600, which agrees
     closely with the theor. value for one arm of the star polymer [mol. weight
     7000]. An analogous polymerization scheme was also developed to prepare graft
     systems.
ΙT
    166983-62-6P
        (tri-functional initiator; architectural control in
        living free radical polymns. with TEMPO derivative functional
        initiators for star and graft polymers)
     166983-62-6 HCAPLUS
RN
     1,3,5-Benzenetricarboxylic acid, tris[2-phenyl-2-[(2,2,6,6-tetramethyl-
CN
     1-piperidinyl)oxy]ethyl] ester (9CI) (CA INDEX NAME)
```

CC 35-3 (Chemistry of Synthetic High Polymers)
 Section cross-reference(s): 36

ST tetramethylpiperidinyloxy deriv functional initiator star polymer; graft polystyrene TEMPO initiator

IT Polymerization catalysts

(TEMPO-derivative initiators; architectural control in living free radical polymns. with TEMPO derivative functional initiators for star and graft polymers)

IT Polymerization

(graft, living free-radical; architectural control in living free radical polymns. with TEMPO derivative functional initiators for star and graft polymers)

IT Polymerization

(radical, living free-radical; architectural control in living free radical polymns. with TEMPO derivative functional initiators for star and graft polymers)

IT Polymers, preparation

IT 1310-58-3, Potassium hydroxide, reactions 1592-20-7,

p-Chloro-methylstyrene 4422-95-1, 1,3,5-Benzenetricarbonyl chloride 81913-53-3

(architectural control in living free radical polymns. with TEMPO derivative functional initiators for star and graft polymers)

IT 161776-41-6P

(architectural control in living free radical polymns. with TEMPO derivative functional initiators for star and graft polymers)

IT 166983-64-8P

(functional copolymer and macroinitiator for preparation of branched polystyrene; architectural control in living free radical polymns. with TEMPO derivative functional initiators for star and graft polymers)

IT 166983-63-7P

(functional graft monomer; architectural control in living free

radical polymns. with TEMPO derivative functional initiators for star and graft polymers)

IT 9003-53-6P, Polystyrene

(star- and comb-branched; architectural control in living free radical polymns. with TEMPO derivative functional initiators for star and graft polymers)

IT 166983-62-6P

(tri-functional initiator; architectural control in living free radical polymns. with TEMPO derivative functional initiators for star and graft polymers)

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                OR 998-40-3/BI)
L3
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L4
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L8
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L9
             50 SEA SSS SAM L9
L10
           7221 SEA SSS FUL L9
L11
L12
             24 SEA ABB=ON PLU=ON L11 AND L2
                SAV L12 KOL030/A
L13
             30 SEA SUB=L11 SSS SAM L5
L14
            694 SEA SUB=L11 SSS FUL L5
                SAV L14 KOL030A/A
T<sub>1</sub>1.5
                STR L9
L16
             50 SEA SUB=L11 SSS SAM L15
L17
                STR L15
L18
             12 SEA SUB=L11 SSS SAM L17
L19
            240 SEA SUB=L11 SSS FUL L17
                SAV L19 KOL030B/A
L20
           6290 SEA ABB=ON PLU=ON L11 NOT (L14 OR L19)
L21
                STR L9
L22
             50 SEA SUB=L11 SSS SAM L21
L23
           2091 SEA SUB=L11 SSS FUL L21
                SAV L23 KOL030C/A
     FILE 'HCAPLUS' ENTERED AT 12:55:27 ON 24 MAR 2008
L24
           262 SEA ABB=ON PLU=ON L14
           144 SEA ABB=ON PLU=ON L19
L25
           789 SEA ABB=ON PLU=ON L23
L26
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6 SEA ABB=ON PLU=ON L12
L27
L28
            1 SEA ABB=ON PLU=ON L27 AND L1
         209 SEA ABB=ON PLU=ON L24(L)PREP/RL
L29
L30
            1 SEA ABB=ON PLU=ON L29 AND CLAY?
           1 SEA ABB=ON PLU=ON L24 AND L25 AND L26
74 SEA ABB=ON PLU=ON L29 AND (PLASTIC? OR POLYMER?)/SC,SX
L31
L32
L33
            60 SEA ABB=ON PLU=ON L32 AND (INITIAT? OR CATALYST? OR
              ACTIVAT?)
            58 SEA ABB=ON PLU=ON L33 NOT L19
L34
   FILE 'REGISTRY' ENTERED AT 13:23:12 ON 24 MAR 2008
L35
              STR L5
L36
            10 SEA SUB=L14 SSS SAM L35
T.3.7
           198 SEA SUB=L14 SSS FUL L35
               SAV L37 KOL030D/A
    FILE 'HCAPLUS' ENTERED AT 13:24:31 ON 24 MAR 2008
      96 SEA ABB=ON PLU=ON L37
L38
            40 SEA ABB=ON PLU=ON L34 AND L38
L39
            32 SEA ABB=ON PLU=ON L26 AND L25
L40
FILE 'REGISTRY' ENTERED AT 13:26:02 ON 24 MAR 2008
              STR L21
L42
              STR L21
           50 SEA SUB=L23 SSS SAM L42
L43
L44
          1677 SEA SUB=L23 SSS FUL L42
               SAV L44 KOL030E/A
    FILE 'HCAPLUS' ENTERED AT 13:32:30 ON 24 MAR 2008
      614 SEA ABB=ON PLU=ON L44
           399 SEA ABB=ON PLU=ON L45(L)PREP/RL
L46
           271 SEA ABB=ON PLU=ON L46 AND (PLASTIC? OR POLYMER?)/SC,SX
L47
           208 SEA ABB=ON PLU=ON L47 AND (INITIAT? OR CATALYST? OR
L48
              ACTIVAT?)
             1 SEA ABB=ON PLU=ON L48 AND L1
L49
L50
             2 SEA ABB=ON PLU=ON L48 AND CLAY?
             4 SEA ABB=ON PLU=ON L48 AND POF/RL
          174 SEA ABB=ON PLU=ON L48 AND CAT/RL
L52
             6 SEA ABB=ON PLU=ON L52 AND MOA/RL
L53
            15 SEA ABB=ON PLU=ON L52 AND (PEP OR PYP)/RL
22 SEA ABB=ON PLU=ON (L49 OR L50 OR L51) OR (L53 OR L54)
L54
L55
L56
           13 SEA ABB=ON PLU=ON L25 AND L46
           35 SEA ABB=ON PLU=ON L55 OR L56
L57
            6 SEA ABB=ON PLU=ON L39 AND ALKOXYAMINE INITIATOR?
           40 SEA ABB=ON PLU=ON L39 OR L58
L59
            5 SEA ABB=ON PLU=ON L57 AND ALKOXYAMINE INITIATOR?
L60
           35 SEA ABB=ON PLU=ON L57 OR L60
26 SEA ABB=ON PLU=ON L45 AND ALKOXYAMINE INITIATOR?
L61
L62
L63
           26 SEA ABB=ON PLU=ON L62 AND (PLASTIC? OR POLYMER?)/SC,SX
L64
           56 SEA ABB=ON PLU=ON L61 OR L63
            6 SEA ABB=ON PLU=ON L38 AND ALKOXYAMINE INITIATOR?
L66
           40 SEA ABB=ON PLU=ON L59 OR L65
           10 SEA ABB=ON PLU=ON L64 AND L66
46 SEA ABB=ON PLU=ON L64 NOT L67
L67
L68
L69 30 SEA ABB=ON PLU=ON L66 NOT
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